

STIC-Biotech/ChemLib

57410

From:
Sent:
To:
Subject:

p

Schnizer, Holly
Wednesday, January 02, 2002 1:21 PM
STIC-Biotech/ChemLib
seq. search request for appl. no. 09/444,281

Please search the commercial and interference databases for

SEQ ID NOs: 27, 35, and 36 and polynucleotides encoding SEQ ID NOs: 27, 35, and 36

Thank you.

Holly Schnizer
AU 1653
CM1-10B05
305-3722
mailbox: CM1-9B01

CRPG

if Contact:
Searcher: Sheppard
Phone: _____
Location: tel: 308-4499
Date Picked Up: _____
Date Completed: 1/8/02
Searcher Prep/Review: _____
Clerical: _____
Online time: _____

TYPE OF SEARCH:
NA Sequences: _____
AA Sequences: _____
Structures: _____
Bibliographic: _____
Litigation: _____
Full text: _____
Patent Family: _____
Other: _____

VENDOR/COST(where applic.)
STN: _____
DIALOG: _____
Questel/Orbit: _____
DRLink: _____
Lexis/Nexis: _____
Sequence Sys.: _____
WWW/Internet: _____
Other (specify): _____

THIS PAGE BLANK (USPTO)

PT conjugates with polyoxyalkylene glycol and fatty acid to reduce
PT toxicity, useful therapeutically, as disinfectants etc.
XX
PS Claim 11; Page 88; 129pp; English.
XX
CC AAY24549 to AAY24615 represent indolicidin analogues of formulae
CC (I)-(VIII) containing up to 25 amino acids (aa): R₁XX₁XB (I), B₁XX₁XB
CC (II), B₁XX₁XX₁XB (III), B₁XX₁XX₁XB₁(AA) (IV), B₁XX₁XX₁XB₁(AA)₁ (V),
CC (VI), LB₁XX₁XX₁XX₁XB₁ (VII), LK₁XX₁XX₁XX₁XB₁ (VIII), B₁XX₁XX₁XB₁ (VIII),
CC (IX) where 2 = P or V; X = hydrophobic residue, preferably W, B = basic aa,
CC preferably R or K; AA = any aa; n = 0 or 1; in (II), at least 1 Z = V;
CC in (VIII) at least 2 X = F or Y. The analogues are used to treat
CC infections caused by bacteria (Gram positive or negative, or anaerobic);
CC fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
CC trematodes) or viruses. Typical of very many pathogens that can be
CC controlled are *Leishmania*, *Trypanosoma*, *Ascaris lumbricoides*, *Pasciella*
CC *hepatica*, *Klebsiella pneumoniae*, *Bordetella pertussis*, *Staphylococcus*
CC *aureus*, *Listeria*, *Clostridium*, *rotavirus* and *papilloma virus*. Compounds
CC derived from the analogues may be used similarly; the compounds may
CC also be prepared from antibiotics or antiarrhythmic agents. The analogues
CC may be used therapeutically or to coat medical devices; also they are
CC useful as surface disinfectants, as additives to shampoo or soaps, as
CC insecticides or herbicides, or as preservatives for foods and technical
CC materials. The analogues are administered by injection, lavage, orally
CC or topically, generally at 0.1-50 mg/kg. These analogues have a broader
CC spectrum of activity than indolicidin and modification as compounds
CC reduces their toxicity.
XX
SQ Sequence 12 AA;
XX

Query Match 100.0%; Score 86; DB 19; Length 12;
Best Local Similarity 100.0%; Pred. No. 3.9e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILRPMWPMWRK 12
| | | | | | | | | | | | | |
Db 1 ILRPMWPMWRK 12

RESULT 2
AAY94496
ID AAY94496 standard; Peptide; 12 AA.
XX
AC AAY94496;
XX
DT 20-SEP-2000 (first entry)
XX
DE MBI-11B7 peptide derived from indolicidin.
XX
DE Cellulose binding domain; CBD; cationic peptide;
KM MBI-11B7, indolicidin; bovine.
XX
OS Bos taurus.
XX
PN WO200031279-A2.
XX
PD 02-JUN-2000.
XX
PE 19-NOV-1999; 99WO-CA01107.
XX
PR 20-NOV-1998; 98US-0109218.
XX
PA (MICR-) MICROLOGIX BIOTECH INC.
XX
PI Burian J, Bartfeld P.
XX
DR WPI: 2000-400086/34.
XX
PT Multi-domain fusion protein expression cassette used for high yield
PT stable production of foreign peptide gene products -
XX
PS Disclosure; Page 24; 73pp; English.

XX
CC A novel method allows the efficient production of cationic peptides in
CC recombinant host cells. The method involves construction of a
CC multi-domain fusion protein expression cassette comprising a promoter and
CC a nucleic acid molecule expressed as an insoluble protein. The inclusion
CC of anionic peptide sequences in the linker sequences neutralises the
CC positive charge of the cationic peptide so that the charge of the
CC fusion protein is controlled. This cassette allows high yield, stable
CC production of the cationic peptide. Cationic peptides such as
CC bovine indolicidin may be used as antimicrobial agents. The present
CC sequence is the MBI-11B7 peptide. MBI-11B7 is a cationic peptide derived
CC from modifications of indolicidin.
XX
SQ Sequence 12 AA;
XX

Query Match 100.0%; Score 86; DB 21; Length 12;
Best Local Similarity 100.0%; Pred. No. 3.9e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILRPMWPMWRK 12
| | | | | | | | | | | | | |
Db 1 ILRPMWPMWRK 12

RESULT 3
AAY91791
ID AAY91791 standard; Peptide; 12 AA.
XX
AC AAY91791;
XX
DT 06-JUN-2000 (first entry)
XX
DE Amino acid sequence of cationic peptide MBI 11B7CN.
XX
KM Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
KM leukaemia; polyoxyalkylene-modified; APO; lymphoma; multiple myeloma;
KM breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
KM multidrug resistance.
XX
OS Synthetic.
XX
PN WO9965506-A2.
XX
PD 23-DEC-1999.
XX
PE 14-JUN-1999; 99WO-CA00552.
XX
PR 12-JUN-1998; 98US-0096541.
XX
PA (MICR-) MICROLOGIX BIOTECH INC.
XX
PI Friedland HD, Krieger TJ, Taylor R, Erle D, Fraser JR, West MHP.
XX
DR WPI: 2000-223549/19.
XX
PT Novel pharmaceutical composition containing optionally activated
PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
XX
PS Claim 1; Page 14; 94pp; English.
XX
CC This sequence represents a cationic peptide amino acid sequence, which
CC can be used in the pharmaceutical composition of the invention. The
CC invention relates to a pharmaceutical composition containing at least one
CC activated polyoxyalkylene (APO)-modified cationic peptide. The
CC modification of peptides with APO increases their activity against tumour
CC cells, including those with a multidrug resistant phenotype. The
CC pharmaceutical composition can be used to treat tumours, specifically
CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
CC cervix, uterus, skin, prostate, liver and colon.
XX
SQ Sequence 12 AA;

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:40:26 ; Search time 53.46 Seconds
(without alignments)
16.627 Million cell updates/sec

Title: US-09-444-281-36
Perfect score: 86
Sequence: 1 ILRPMWPMWRRK 12

Scoring table: BIOSUM62
Gapop 10.0, Gapext 0.5

Searched: 522463 seqs, 74073290 residues

Total number of hits satisfying chosen parameters: 522463

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 08
Maximum Match 1008
Listing first 45 summaries

Database :

A_Geneseq.1101.*
1: /SIDSR/gcgdata/geneseq/geneseq/AA1980.DAT.*
2: /SIDSR/gcgdata/geneseq/geneseq/AA1981.DAT.*
3: /SIDSR/gcgdata/geneseq/geneseq/AA1982.DAT.*
4: /SIDSR/gcgdata/geneseq/geneseq/AA1983.DAT.*
5: /SIDSR/gcgdata/geneseq/geneseq/AA1984.DAT.*
6: /SIDSR/gcgdata/geneseq/geneseq/AA1985.DAT.*
7: /SIDSR/gcgdata/geneseq/geneseq/AA1986.DAT.*
8: /SIDSR/gcgdata/geneseq/geneseq/AA1987.DAT.*
9: /SIDSR/gcgdata/geneseq/geneseq/AA1988.DAT.*
10: /SIDSR/gcgdata/geneseq/geneseq/AA1989.DAT.*
11: /SIDSR/gcgdata/geneseq/geneseq/AA1990.DAT.*
12: /SIDSR/gcgdata/geneseq/geneseq/AA1991.DAT.*
13: /SIDSR/gcgdata/geneseq/geneseq/AA1992.DAT.*
14: /SIDSR/gcgdata/geneseq/geneseq/AA1993.DAT.*
15: /SIDSR/gcgdata/geneseq/geneseq/AA1994.DAT.*
16: /SIDSR/gcgdata/geneseq/geneseq/AA1995.DAT.*
17: /SIDSR/gcgdata/geneseq/geneseq/AA1996.DAT.*
18: /SIDSR/gcgdata/geneseq/geneseq/AA1997.DAT.*
19: /SIDSR/gcgdata/geneseq/geneseq/AA1998.DAT.*
20: /SIDSR/gcgdata/geneseq/geneseq/AA1999.DAT.*
21: /SIDSR/gcgdata/geneseq/geneseq/AA2000.DAT.*
22: /SIDSR/gcgdata/geneseq/geneseq/AA2001.DAT.*

pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	86	100.0	12	19	AAV24550
2	86	100.0	12	21	AAV94496
3	86	100.0	12	21	AAV91791
4	86	100.0	20	19	AAV24553
5	86	100.0	20	21	AAV91797
6	86	100.0	21	19	AAV24552
7	86	100.0	21	19	AAV24554
8	86	100.0	21	19	AAV66376
9	86	100.0	21	21	AAV91796
10	86	100.0	21	21	AAV91798
11	86	100.0	27	19	AAV66363

12	86	100.0	28	21	AAV91800
13	83	96.5	12	19	AAV24567
14	82	96.5	12	21	AAV91788
15	82	95.3	12	19	AAV24594
16	82	95.3	12	19	AAV66364
17	82	95.3	12	21	AAV91817
18	82	95.3	12	21	AAV91841
19	81	94.2	12	19	AAV24605
20	81	94.2	12	19	AAV24595
21	81	94.2	12	21	AAV91842
22	81	94.2	12	21	AAV91852
23	80	93.0	12	19	AAV24596
24	80	93.0	12	19	AAV24603
25	80	93.0	12	19	AAV24604
26	80	93.0	12	21	AAV91843
27	80	93.0	12	21	AAV91850
28	80	93.0	12	21	AAV91851
29	78	90.7	12	19	AAV24598
30	78	90.7	12	19	AAV24601
31	78	90.7	12	19	AAV66361
32	78	90.7	12	21	AAV91785
33	78	90.7	12	21	AAV91845
34	78	90.7	12	21	AAV91848
35	78	90.7	13	19	AAV24565
36	78	90.7	13	21	AAV24586
37	77	89.5	12	19	AAV24586
38	77	89.5	12	21	AAV91786
39	75	87.2	11	19	AAV24569
40	75	87.2	11	21	AAV91790
41	75	87.2	12	19	AAV24580
42	75	87.2	12	21	AAV91804
43	75	87.2	13	18	AAV12873
44	75	87.2	13	18	AAV12895
45	75	87.2	13	18	AAV12896

ALIGNMENTS

RESULT 1
ID AAV24550 standard; peptide; 12 AA.
AC AAV24550;
XX
XX 18-AUG-1999 (first entry)
DT
XX
DE Indolicidin analogue #2.
KW Indolicidin; bacterial infection; photo-oxidised solubiliser;
KW antimicrobial; antibiotic; antidiarrhythmic; surface disinfectant;
KW additive; shampoo; soap; insecticide; herbicide; preservative;
KW food; technical material.
XX
XX Synthetic.
OS
XX
XX WO9807745-A2.
XX
XX 26-FEB-1998.
PD
XX
XX 21-AUG-1997; 97WO-US14779.
PF
XX
XX 13-JAN-1997; 97US-0034949.
PR
XX 21-AUG-1996; 96US-0024754.
XX
XX (MICR-) MICROLOGIX BIOTECH INC.
XX
XX Effle D, Fraser JR, Krieger TU, Taylor R, West MH;
WPI; 1998-169090/15.
XX
XX New indolicidin analogues with antimicrobial activity and related
PT nucleic acid - vectors, transformed cells and antibodies, also

Query Match 100.0%; Score 86; DB 21; Length 12;
Best Local Similarity 100.0%; Pred. No. 3.9e-06;
Matches 12: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRPMWPMRRK 12
| | | | | | | | | | | |
DB 1 ILRPMWPMRRK 12

RESULT 4
AAV24553
ID AAV24553 standard; peptide: 20 AA.
XX
AC AAV24553;
XX
DT 18-AUG-1999 (first entry)
XX
DE Indolicidin analogue #5.
XX
KW Indolicidin; bacterial infection; photo-oxidised solubiliser;
KM antimicrobial; antibiotic; antitarrhythmic; surface disinfectant;
KW additive; shampoo; soap; insecticide; herbicide; preservative;
KW food; technical material.
XX
OS Synthetic.
XX
PN WO9807745-A2.
XX
PD 26-FEB-1998.
XX
PE 21-AUG-1997; 97WO-US14779.
XX
PF 13-JAN-1997; 97US-0034949.
PR 21-AUG-1996; 96US-0024754.
XX
PA (MICR-) MICROLOGIX BIOTECH INC.
XX
PI Erfle D, Fraser JR, Krieger TJ, Taylor R, West MH;
DR WPI; 1998-169090/15.
XX
PT New indolicidin analogues with antimicrobial activity and related
PT nucleic acid - vectors, transformed cells and antibodies, also
PT conjugates with polyoxyalkylene glycol and fatty acid to reduce
PT toxicity, useful therapeutically, as disinfectants etc.
XX
PS Claim 11; Page 88; 129pp; English.
XX
CC AAV24549 to AAV24615 represent indolicidin analogues of formulae
CC (I)-(VIII) containing up to 25 amino acids (aa): RZXZXZXB (I), BXZXZXZB
CC (II), BBBZXZXZXB (III), BXZXZXZBBn(A)nmlBBAGS (IV), BXZXZXZBB(A)nM
CC (V), LBBnZXZXZnXRK (VI), LKnZXZXZXRK (VII) and BBXZXZXZBBB (VIII).
CC Where Z = P or V; X = hydrophobic residue, preferably W; B = basic aa,
CC preferably R or K; AA = any aa; n = 0 or 1; in (II), at least 1 Z = V;
CC in (VIII) at least 2 X = F or Y. The analogues are used to treat
CC infections caused by bacteria (Gram positive or negative, or anaerobic);
CC fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
CC trematodes) or viruses. Typical of very many pathogens that can be
CC controlled are Leishmania, Trypanosoma, Ascaris lumbricoides, Fasciola
CC hepatica, Klebsiella pneumoniae, Bordetella pertussis, Staphylococcus
CC aureus, Listeria, Clostridium, rotavirus and papilloma virus. Compounds
CC derived from the analogues may be used similarly; the compounds may
CC also be prepared from antibiotics or antitarrhythmic agents. The analogues
CC may be used therapeutically or to coat medical devices; also they are
CC useful as surface disinfectants, as additives to shampoo or soaps, as
CC insecticides or herbicides, or as preservatives for foods and technical
CC materials. The analogues are administered by injection, lavage, orally
CC or topically, generally at 0.1-50 mg/Kg. These analogues have a broader
CC spectrum of activity than indolicidin and modification as compounds
CC reduces their toxicity.
XX

SQ Sequence 20 AA;

Query Match 100.0%; Score 86; DB 19; Length 20;
Best Local Similarity 100.0%; Pred. No. 6.5e-06;
Matches 12: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRPMWPMRRK 12
| | | | | | | | | | | |
DB 1 ILRPMWPMRRK 12

RESULT 5
AAV91797
ID AAV91797 standard; peptide: 20 AA.
XX
AC AAV91797;
XX
DT 06-JUN-2000 (first entry)
XX
DE Amino acid sequence of cationic peptide MBI 11B17CN.
XX
KW Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
KM leukaemia; polyoxyalkylene-modified; APO; lymphoma; multiple myeloma;
KW breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
KW multidrug resistance.
XX
OS Synthetic.
XX
PN WO9965506-A2.
XX
PD 23-DEC-1999.
XX
PE 14-JUN-1999; 99WO-CA00552.
XX
PR 12-JUN-1998; 98US-0096541.
XX
PA (MICR-) MICROLOGIX BIOTECH INC.
XX
PI Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP;
DR WPI; 2000-223549/19.
XX
PT Novel pharmaceutical composition containing optionally activated
PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
XX
PS Disclosure; Page 15; 94pp; English.
XX
CC This sequence represents a cationic peptide amino acid sequence, which
CC can be used in the pharmaceutical composition of the invention. The
CC invention relates to a pharmaceutical composition containing at least one
CC activated polyoxyalkylene (APO)-modified cationic peptide. The
CC modification of peptides with APO increases their activity against tumour
CC cells, including those with a multidrug resistant phenotype. The
CC pharmaceutical composition can be used to treat tumours, specifically
CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
CC cervix, uterus, skin, prostate, liver and colon.
XX
SQ Sequence 20 AA;

Query Match 100.0%; Score 86; DB 21; Length 20;
Best Local Similarity 100.0%; Pred. No. 6.5e-06;
Matches 12: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRPMWPMRRK 12
| | | | | | | | | | | |
DB 1 ILRPMWPMRRK 12

RESULT 6
AAV24552

ID AAY24552 standard; peptide; 21 AA.
 AC AAY24552;
 XX
 DT 18-AUG-1999 (first entry)
 XX
 DE Indolicidin analogue #4.
 XX
 KM Indolicidin; bacterial infection; photo-oxidised solubiliser;
 KM antimicrobial; antibiotic; antitarrhythmic; surface disinfectant;
 KW additive; Shampoo; soap; insecticide; herbicide; preservative;
 KM food; technical material.
 OS Synthetic.
 PN W09807745-A2.
 PN
 PD 26-FEB-1998.
 PD
 XX 21-AUG-1997; 97WO-US14779.
 XX
 PF 13-JAN-1997; 97US-0034949.
 PR 21-AUG-1996; 96US-0024734.
 XX
 PA (MCR-) MICROLOGIX BIOTECH INC.
 PA
 PI Ernie D, Fraser JR, Krieger TJ, Taylor R, West MH;
 PI
 DR WPI: 1998-169090/15.
 XX
 PT New indolicidin analogues with antimicrobial activity and related
 PT nucleic acid - vectors, transformed cells and antibodies, also
 PT conjugates with polyoxyalkylene glycol and fatty acid to reduce
 PT toxicity, useful therapeutically, as disinfectants etc.
 XX
 PS Claim 11; Page 88; 129pp; English.
 PS
 XX AAY24549 to AAY24615 represent indolicidin analogues of formulae
 CC (I)-(VIII) containing up to 25 amino acids (aa): R₁XXXXXB (I), B₁XXXXXB
 CC (II), B₁B₂XXXXXB (III), B₁X₁X₂X₃B₁B₂B₃(AA)nmLB₁B₂B₃GS (IV), B₁X₁X₂X₃B₁(AA)n
 CC (V), LB₁B₂B₃nmXXXXXB (VI), LK₁X₁X₂X₃RRK (VII) and B₁X₁X₂X₃B₁B₂B₃ (VIII).
 CC Where Z = P or V; X = hydrophobic residue, preferably W; B = basic aa,
 CC preferably R or K; AA = any aa; n = 0 or 1, in (II), at least 1 Z = V;
 CC in (VIII) at least 2 X = F or Y. The analogues are used to treat
 CC infections caused by bacteria (gram positive or negative, or anaerobic);
 CC fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
 CC trematodes) or viruses. Typical of very many pathogens that can be
 CC controlled are Leishmania, Trypanosoma, Ascaris lumbricoides, Fasciola
 CC hepatica, Klebsiella pneumoniae, Bordetella pertussis, Staphylococcus
 CC aureus, Listeria, Clostridium, rotavirus and papilloma virus. Compounds
 CC derived from the analogues may be used similarly: the compounds may
 CC also be prepared from antibiotics or antitarrhythmic agents. The analogues
 CC may be used therapeutically or to coat medical devices; also they are
 CC useful as surface disinfectants, as additives to shampoo or soaps, as
 CC insecticides or herbicides, or as preservatives for foods and technical
 CC materials. The analogues are administered by injection, lavage, orally
 CC or topically, generally at 0.1-50 mg/kg. These analogues have a broader
 CC spectrum of activity than indolicidin and modification as compounds
 CC reduces their toxicity.
 XX
 XX Sequence 21 AA;
 XX

ID AAY24554 standard; peptide; 21 AA.
 AC AAY24554;
 XX
 DT 18-AUG-1999 (first entry)
 XX
 XX Indolicidin analogue #6.
 DE
 KW Indolicidin; bacterial infection; photo-oxidised solubiliser;
 KW antimicrobial; antibiotic; antiarrhythmic; surface disinfectant;
 KW additive; shampoo; soap; insecticide; herbicide; preservative;
 KW food; technical material.
 XX
 OS Synthetic.
 PN WO9807745-A2.
 XX
 PD 26-FEB-1998.
 XX
 PE 21-AUG-1997; 97WO-US14779.
 XX
 PR 13-JAN-1997; 97US-0034949.
 PR 21-AUG-1996; 96US-0024754.
 PA (MICR-) MICROLOGIX BIOTECH INC.
 PI Erle D, Fraser JR, Krieger TJ, Taylor R, West MH;
 XX
 DR WPI; 1998-169090/15.
 XX
 PT New indolicidin analogues with antimicrobial activity and related
 PT nucleic acid - vectors; transformed cells and antibodies, also
 PT conjugates with polyoxalkylene glycol and fatty acid to reduce
 PT toxicity, useful therapeutically, as disinfectants etc.
 XX
 XX Claim 11; Page 88; 129pp; English.
 PS
 XX
 AA AAY24549 to AAY24615 represent indolicidin analogues of formulae
 (I)-(VIII) containing up to 25 amino acids (aa): RX_2XX_2XB (I), BX_2XX_2XB
 (II), BBX_2XX_2XB (III), $BX_2XX_2XBBn(AA)nMILBAGS$ (IV), $BX_2XX_2XB(AA)nM$
 (V), $LBnX_2XX_2XNRK$ (VI), $LKnX_2XX_2XNRK$ (VII) and BBX_2XX_2XBBB (VIII).
 Where Z = P or V; X = hydrophobic residue, preferably W; B = basic aa,
 preferably R or K; AA = any aa; n = 0 or 1; in (II), at least 1 Z = V;
 in (VIII) at least 2 X = F or Y. The analogues are used to treat
 infections caused by bacteria (Gram positive or negative, or anaerobic);
 fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
 trematodes) or viruses. Typical of very many pathogens that can be
 controlled are Leishmania, Trypanosoma, Ascaris lumbricoides, Fasciola
 hepatica, Klebsiella pneumoniae, Bordetella pertussis, Staphylococcus
 aureus, Listeria, Clostridium, rotavirus and papilloma virus. Compounds
 derived from the analogues may be used similarly: the compounds may
 also be prepared from antibiotics or antiarrhythmic agents. The analogues
 may be used therapeutically or to coat medical devices; also they are
 useful as surface disinfectants, as additives to shampoo or soaps, as
 insecticides or herbicides, or as preservatives for foods and technical
 materials. The analogues are administered by injection, lavage, orally
 or topically, generally at 0.1-50 mg/kg. These analogues have a broader
 spectrum of activity than indolicidin and modification as compounds
 reduces their toxicity.
 XX
 Sequence 21 AA;
 50

RESULT 8
 AAM6376 standard; peptide: 21 AA.
 AC AAM6376;
 DT 12-JAN-1999 (first entry)
 DE Cationic peptide of claim 15 #3.
 KM Indolicidin analogue; resistance; cationic peptide; antibiotic;
 KM bacterial infection; tolerance; antibacterial; microorganism;
 KM bacteria; fungus; parasite; virus.
 OS Synthetic.
 PN WO9840401-A2.
 PD 17-SEP-1998.
 PF 10-MAR-1998; 98WO-CA00190.
 PR 25-FEB-1998; 98US-0030619.
 PR 10-MAR-1997; 97US-0040649.
 PR 20-AUG-1997; 97US-0915314.
 PR 26-SEP-1997; 97US-0060099.
 PS (MICR-) MICROLOGIX BIOTECH INC.
 PI Fraser JR, McNicol PJ, West MHP;
 DR WPI: 1998-520800/44.
 XX
 XX WPI: 1998-520800/44.
 DR
 XX
 XX New indolicidin peptide analogues - useful for, e.g. enhancing
 PT activity of antibiotic or overcoming tolerance, acquired resistance
 PT or inherent resistance of microorganisms
 PS
 PS Claim 15; Page 93; 105pp; English.
 CC The present sequence represents a specifically claimed cationic peptide
 CC from the present invention. The present invention describes compositions
 CC and methods for treating infection, especially bacterial infections. The
 CC compositions and methods use cationic peptides in combination with an
 CC antibiotic agent which are then administered to a patient to enhance the
 CC activity of the antibiotic agent, to overcome: (a) tolerance; (b)
 CC acquired resistance; and (c) inherent resistance. The combinations of
 CC antibiotics and cationic peptides can provide synergistic activity
 CC against a microorganism that is tolerant, inherently resistant, or has
 CC acquired resistance to an antibiotic agent. They can be used for killing
 CC e.g. bacteria, fungi, parasites and viruses.
 CC
 SQ Sequence 21 AA:

Query Match 100.0%; Score 86; DB 19; Length 21;
 Best Local Similarity 100.0%; Pred. No. 6.8e-06;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRWPMWPMRRK 12
 DB 1 ILRWPMWPMRRK 12

RESULT 9
 AAY91796 standard; peptide: 21 AA.
 AC AAY91796;
 DT 06-JUN-2000 (first entry)
 DE Amino acid sequence of cationic peptide MBI 11B16CN.

KM Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
 KM leukaemia; polyoxalkylene-modified; APO; lymphoma; multiple myeloma;
 KM breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
 KM multidrug resistance.
 OS Synthetic.
 PN WO9965506-A2.
 PD 23-DEC-1999.
 PF 14-JUN-1999; 99WO-CA00552.
 PR 12-JUN-1998; 98US-0096541.
 PS (MICR-) MICROLOGIX BIOTECH INC.
 PI Friedland HD, Krieger TJ, Taylor R, Effie D, Fraser JR, West MHP;
 DR WPI: 2000-223549/19.
 XX
 XX
 XX Novel pharmaceutical composition containing optionally activated
 PT polyoxalkylene-modified cationic peptides, useful for treating tumours
 PT
 PS Disclosure: Page 15; 94pp; English.
 CC This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.
 CC
 SQ Sequence 21 AA:

Query Match 100.0%; Score 86; DB 21; Length 21;
 Best Local Similarity 100.0%; Pred. No. 6.8e-06;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRWPMWPMRRK 12
 DB 1 ILRWPMWPMRRK 12

RESULT 10
 AAY91798 standard; peptide: 21 AA.
 AC AAY91798;
 DT 06-JUN-2000 (first entry)
 DE Amino acid sequence of cationic peptide MBI 11B18CN.
 KM Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
 KM leukaemia; polyoxalkylene-modified; APO; lymphoma; multiple myeloma;
 KM breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
 KM multidrug resistance.
 OS Synthetic.
 PN WO9965506-A2.
 PD 23-DEC-1999.
 PF 14-JUN-1999; 99WO-CA00552.
 PR 12-JUN-1998; 98US-0096541.

XX (MICR-) MICROLOGIX BIOTECH INC.
 XX Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP;
 PI WPI; 2000-223549/19.
 XX Novel pharmaceutical composition containing optionally activated
 PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
 CC
 PS Disclosure; Page 15; 94pp; English.
 XX
 CC This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxyalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.
 CC
 SQ Sequence 21 AA;

Query Match 100.0%; Score 86; DB 21; Length 21;
 Best Local Similarity 100.0%; Pred. No. 6.8e-06;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRMPWMPWRRK 12
 |||||
 DB 1 ILRMPWMPWRRK 12

RESULT 11

AAM66363
 ID AAM66363 standard; peptide; 27 AA.

AC AAM66363;

DT 12-JAN-1999 (first entry)

DE Indolicidin analogue MBI 11B20.

KW Indolicidin analogue; resistance; cationic peptide; antibiotic;

KW bacterial infection; tolerance; antibacterial; microorganism;

OS Bos taurus.

OS Synthetic.

PN WO9840401-A2.

PD 17-SEP-1998.

PF 10-MAR-1998; 98WO-CA00190.

PR 25-FEB-1998; 98US-0030619.

PR 10-MAR-1997; 97US-0040649.

PR 20-AUG-1997; 97US-0915314.

PR 26-SEP-1997; 97US-0060099.

PA (MICR-) MICROLOGIX BIOTECH INC.

PI Fraser JR, McNicol PJ, West MHP;

DR WPI; 1998-520800/44.

XX New indolicidin peptide analogues - useful for, e.g. enhancing

PT activity of antibiotic or overcoming tolerance, acquired resistance

XX or inherent resistance of microorganisms

PS Claim 1; Page 91; 105pp; English.

XX The present sequence represents an indolicidin analogue. The present
 CC invention describes compositions and methods for treating infection,
 CC especially bacterial infections. The compositions and methods use
 CC cationic peptides in combination with an antibiotic agent which are
 CC then administered to a patient to enhance the activity of the antibiotic
 CC agent, to overcome: (a) tolerance; (b) acquired resistance; and (c)
 CC inherent resistance. The combinations of antibiotics and cationic
 CC peptides can provide synergistic activity against a microorganism that
 CC is tolerant, inherently resistant, or has acquired resistance to an
 CC antibiotic agent. They can be used for killing e.g. bacteria, fungi,
 CC parasites and viruses.
 CC
 SQ Sequence 27 AA;

Query Match 100.0%; Score 86; DB 19; Length 27;
 Best Local Similarity 100.0%; Pred. No. 8.8e-06;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRMPWMPWRRK 12
 |||||
 DB 1 ILRMPWMPWRRK 12

RESULT 12

AA91800
 ID AA91800 standard; Peptide; 28 AA.

AC AA91800;

DT 06-JUN-2000 (first entry)

DE Amino acid sequence of cationic peptide MBI 11B20CN.

KW Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;

KW leukaemia; polyoxyalkylene-modified; APO; lymphoma; multiple myeloma;

KW breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;

OS Synthetic.

PN WO965506-A2.

PD 23-DEC-1999.

PF 14-JUN-1999; 99WO-CA00552.

PR 12-JUN-1998; 98US-0096541.

PA (MICR-) MICROLOGIX BIOTECH INC.

PI Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP;

DR WPI; 2000-223549/19.

XX Novel pharmaceutical composition containing optionally activated

PT polyoxyalkylene-modified cationic peptides, useful for treating tumours

PS Claim 1; Page 15; 94pp; English.

XX This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxyalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.

SQ Sequence 28 AA:

Query Match 100.0%; Score 86; DB 21; Length 28;
 Best Local Similarity 100.0%; Pred. No. 9.1e-06;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRPMWPMRRK 12
 ||:|||||
 DB 1 ILRPMWPMRRK 12

RESULT 13

AAV24567
 ID AAV24567 standard; peptide: 12 AA.

AC AAV24567;

DT 18-AUG-1999 (first entry)

DE Indolicidin analogue #19.

KW Indolicidin; bacterial infection; photo-oxidised solubiliser;
 KW antimicrobial; antibiotic; antiarrhythmic; surface disinfectant;
 KW additive; shampoo; soap; insecticide; herbicide; preservative;
 KW food; technical material.

OS Synthetic.

PN WO9807745-A2.

PD 26-FEB-1998.

PF 21-AUG-1997; 97WO-US14779.

PR 13-JAN-1997; 97US-0034949.

PR 21-AUG-1996; 96US-0024754.

PA (MICR-) MICROLOGIX BIOTECH INC.

PI Erle D, Fraser JR, Krieger TJ, Taylor R, West MH;

DR WPI: 1998-169090/15.

PT New indolicidin analogues with antimicrobial activity and related
 PT nucleic acid vectors, transformed cells and antibodies, also
 PT conjugates with polyoxyalkylene glycol and fatty acid to reduce
 PT toxicity, useful therapeutically, as disinfectants etc.

PS Claim 12; Page 89; 129pp; English.

XX AA24549 to AAV24615 represent indolicidin analogues of formulae
 CC (I)-(VIII) containing up to 25 amino acids (aa): RXXXXZB (I), BXXXXZB
 CC (II), BBXXXXZB (III), BXXXXZBB(AA)nmILBAGS (IV), BXXXXZB(AA)nm
 CC (V), LBnmZnXXZB (VI), LKXXXXZB (VII) and BXXXXZBB (VIII).
 CC Where Z = P or V; X = hydrophobic residue, preferably W; B = basic aa,
 CC preferably R or K; AA = any aa; n = 0 or 1; in (II), at least 1 Z = V;
 CC in (VIII) at least 2 X = F or Y. The analogues are used to treat
 CC infections caused by bacteria (Gram positive or negative, or anaerobic);
 CC fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
 CC trematodes) or viruses. Typical of very many pathogens that can be
 CC controlled are Leishmania, Trypanosoma, Ascaris lumbricoides, Fasciola
 CC hepatica, Klebsiella pneumoniae, Bordetella pertussis, Staphylococcus
 CC aureus, Listeria, Clostridium, rotavirus and papilloma virus. Compounds
 CC derived from the analogues may be used similarly: the compounds may
 CC also be prepared from antibiotics or antiarrhythmic agents. The analogues
 CC may be used therapeutically or to coat medical devices; also they are
 CC useful as surface disinfectants, as additives to shampoo or soaps, as
 CC insecticides or herbicides, or as preservatives for foods and technical
 CC materials. The analogues are administered by injection, lavage, orally
 CC or topically, generally at 0.1-50 mg/kg. These analogues have a broader
 CC spectrum of activity than indolicidin and modification as compounds
 CC reduces their toxicity.

XX SQ Sequence 12 AA:

Query Match 96.5%; Score 83; DB 19; Length 12;
 Best Local Similarity 91.7%; Pred. No. 1e-05;
 Matches 11; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRPMWPMRRK 12
 ||:|||||
 DB 1 ILKPMWPMRRK 12

RESULT 14

AAV91788
 ID AAV91788 standard; peptide: 12 AA.

AC AAV91788;

DT 06-JUN-2000 (first entry)

DE Amino acid sequence of cationic peptide MBI 11B3CN.

KW Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
 KW leukaemia; polyoxyalkylene-modified; APO; lymphoma; multiple myeloma;
 KW breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
 KW multidrug resistance.

OS Synthetic.

PN WO965506-A2.

PD 23-DEC-1999.

PF 14-JUN-1999; 99WO-CA00552.

PR 12-JUN-1998; 98US-0096541.

PA (MICR-) MICROLOGIX BIOTECH INC.

PI Friedland HD, Krieger TJ, Taylor R, Erle D, Fraser JR, West MHP;

DR WPI: 2000-223549/19.

PT Novel pharmaceutical composition containing optionally activated
 PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
 PT .
 PS Disclosure: Page 14; 94pp; English.

XX This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxyalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.

SQ Sequence 12 AA:

Query Match 96.5%; Score 83; DB 21; Length 12;
 Best Local Similarity 91.7%; Pred. No. 1e-05;
 Matches 11; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRPMWPMRRK 12
 ||:|||||
 DB 1 ILKPMWPMRRK 12

RESULT 15

Search completed: January 4, 2002, 08:40:26
Job time: 109 sec

AAV24594
ID AAV24594 standard; peptide; 12 AA.
XX
AC AAV24594;
XX
DT 18-AUG-1999 (first entry)
XX
DE Indolicidin analogue #46.
XX
KW Indolicidin; bacterial infection; photo-oxidised solubiliser;
KW antimicrobial; antibiotic; antiarrhythmic; surface disinfectant;
KW additive; shampoo; soap; insecticide; herbicide; preservative;
KW food; technical material.
XX
OS Synthetic.
XX
PN W09807745-A2.
XX
PD 26-FEB-1998.
XX
PF 21-AUG-1997; 97WO-US14779.
XX
PR 13-JAN-1997; 97US-0034949.
PR 21-AUG-1996; 96US-0024754.
XX
PA (MICR-) MICROLOGIX BIOTECH INC.
PI Erfle D, Frazer JR, Krieger TJ, Taylor R, West MH;
XX
DR WPI; 1998-169090/15.
XX
PT New indolicidin analogues with antimicrobial activity and related
PT nucleic acid - vectors, transformed cells and antibodies, also
PT conjugates with polyoxyalkylene glycol and fatty acid to reduce
PT toxicity, useful therapeutically, as disinfectants etc.
XX
PS Claim 14; Page 89; 129pp; English.
XX
XX AAV24549 to AAV24615 represent indolicidin analogues of formulae
CC (I)-(VII) containing up to 25 amino acids (aa): RXZXXXB (I), BXZXXXB
CC (II), BBXZXXZXB (III), BXZXXZBBn(AA)nLBAGS (IV), BXZXXXB(AA)nM
CC (V), LBnXZnXnXnXRK (VI), LKnXZnXnXRK (VII) and BBXZXXZBBB (VIII).
CC Where Z = P or V; X = hydrophobic residue, preferably W; B = basic aa,
CC preferably R or K; AA = any aa; n = 0 or 1; In (II), at least 1 Z = V;
CC in (VIII) at least 2 X = F or Y. The analogues are used to treat
CC infections caused by bacteria (Gram positive or negative, or anaerobic);
CC fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
CC trematodes) or viruses. Typical of very many pathogens that can be
CC controlled are Leishmania, Trypanosoma, Ascaris lumbricoides, Fasciola
CC hepatica, Klebsiella pneumoniae, Bordetella pertussis, Staphylococcus
CC aureus, Listeria, Clostridium, rotavirus and papilloma virus. Compounds
CC derived from the analogues may be used similarly: the compounds may
CC also be prepared from antibiotics or antiarrhythmic agents. The analogues
CC may be used therapeutically or to coat medical devices: also they are
CC useful as surface disinfectants, as additives to shampoo or soaps, as
CC insecticides or herbicides, or as preservatives for foods and technical
CC materials. The analogues are administered by injection, lavage, orally
CC or topically, generally at 0.4-50 mg/kg. These analogues have a broader
CC spectrum of activity than indolicidin and modification as compounds
CC reduces their toxicity.
XX
SQ Sequence 12 AA;
XX

Query Match 95.3%; Score 82; DB 19; Length 12;
Best Local Similarity 100.0%; Pred. No. 1.4e-05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 LRPWMPWRRK 12
| | | | | | | | | | | | | |
DB 2 LRWPWPWRRK 12

GenCore version 4.5
Copyright: (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:40:57 ; Search time 24.75 Seconds
(without alignments)
10.911 Million cell updates/sec

Title: US-09-444-281-36
Perfect score: 86
Sequence: 1 ILRPMWPMRRK 12

Scoring table: BLOSUM62
Gapop 10.0, Gapext 0.5

Searched: 212252 seqs, 22503292 residues

Total number of hits satisfying chosen parameters: 212252

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents-AA:*
1: /cgn2_6/ptodata/2/1aa/5A.COMB.pep:*
2: /cgn2_6/ptodata/2/1aa/5B.COMB.pep:*
3: /cgn2_6/ptodata/2/1aa/6A.COMB.pep:*
4: /cgn2_6/ptodata/2/1aa/6B.COMB.pep:*
5: /cgn2_6/ptodata/2/1aa/PCYUS.COMB.pep:*
6: /cgn2_6/ptodata/2/1aa/Backfill1.pep:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	86	100.0	12	US-08-915-314-42	Sequence 42, Appl
2	86	100.0	20	US-08-915-314-47	Sequence 47, Appl
3	86	100.0	21	US-08-915-314-46	Sequence 46, Appl
4	86	100.0	21	US-08-915-314-48	Sequence 48, Appl
5	83	96.5	12	US-08-915-314-40	Sequence 40, Appl
6	82	95.3	12	US-08-915-314-76	Sequence 76, Appl
7	81	94.2	12	US-08-915-314-77	Sequence 77, Appl
8	81	94.2	12	US-08-915-314-78	Sequence 78, Appl
9	80	93.0	12	US-08-915-314-85	Sequence 85, Appl
10	80	93.0	12	US-08-915-314-86	Sequence 86, Appl
11	80	93.0	12	US-08-915-314-80	Sequence 80, Appl
12	78	90.7	12	US-08-915-314-83	Sequence 83, Appl
13	78	90.7	12	US-08-915-314-88	Sequence 88, Appl
14	77	89.5	12	US-08-915-314-38	Sequence 38, Appl
15	77	89.5	12	US-08-915-314-69	Sequence 69, Appl
16	75	87.2	11	US-08-915-314-41	Sequence 41, Appl
17	75	87.2	12	US-08-915-314-52	Sequence 52, Appl
18	75	87.2	13	US-08-915-314-25	Sequence 25, Appl
19	75	87.2	13	US-08-915-314-30	Sequence 30, Appl
20	75	87.2	13	US-08-915-314-51	Sequence 51, Appl
21	75	87.2	13	US-08-915-314-62	Sequence 62, Appl
22	75	87.2	13	US-08-915-314-63	Sequence 63, Appl
23	75	87.2	13	US-08-915-314-64	Sequence 64, Appl
24	75	87.2	13	US-08-702-054B-33	Sequence 33, Appl
25	75	87.2	13	US-08-702-054B-34	Sequence 34, Appl
26	75	87.2	13	US-08-702-054B-35	Sequence 35, Appl
27	75	87.2	13	US-09-042-071-36	Sequence 36, Appl

28	75	87.2	14	US-08-915-314-57	Sequence 57, Appl
29	75	87.2	16	US-08-702-054B-2	Sequence 2, Appl
30	75	87.2	21	US-08-915-314-54	Sequence 54, Appl
31	73	84.9	9	US-08-915-314-90	Sequence 90, Appl
32	73	84.9	11	US-08-915-314-44	Sequence 44, Appl
33	73	84.9	16	US-08-702-054B-38	Sequence 38, Appl
34	72	83.7	12	US-08-915-314-79	Sequence 79, Appl
35	72	83.7	12	US-08-915-314-81	Sequence 81, Appl
36	72	83.7	12	US-08-915-314-82	Sequence 82, Appl
37	72	83.7	12	US-08-915-314-84	Sequence 84, Appl
38	70	81.4	9	US-09-076-227-5	Sequence 5, Appl
39	70	81.4	10	US-09-076-227-4	Sequence 4, Appl
40	70	81.4	11	US-08-702-054B-9	Sequence 9, Appl
41	70	81.4	11	US-09-076-227-3	Sequence 3, Appl
42	70	81.4	12	US-08-915-314-39	Sequence 39, Appl
43	70	81.4	12	US-08-915-314-74	Sequence 74, Appl
44	70	81.4	12	US-08-702-054B-5	Sequence 5, Appl
45	70	81.4	12	US-09-076-227-2	Sequence 2, Appl

ALIGNMENTS

RESULT 1
US-08-915-314-42
; Sequence 42, Application US/08915314
; Patent No. 6180604
; GENERAL INFORMATION:
; APPLICANT: Fraser, Janet R.
; APPLICANT: West, Michael H.P.
; APPLICANT: Krieger, Timothy J.
; APPLICANT: Taylor, Robert
; APPLICANT: Erlic, Douglas
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
; TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
; NUMBER OF SEQUENCES: 90
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SEED and BERRY LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: Washington
; COUNTRY: USA
; ZIP: 98104
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/915,314
; FILING DATE: 20-AUG-1997
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: No. 6180604lenburg Ph.D., Carol
; REGISTRATION NUMBER: 39,317
; REFERENCE/DOCKET NUMBER: 660081.405
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (206) 622-4900
; TELEFAX: (206) 682-6031
; INFORMATION FOR SEQ ID NO: 42:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 12 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; US-08-915-314-42

Query Match 100.0%; Score 86; DB 4; Length 12;
Best Local Similarity 100.0%; Pred No. 1.3e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ILRPMWPMRRK 12

DB 1 ILRWPMPWRRK 12

RESULT 2

US-08-915-314-47
Sequence 47, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 47:
SEQUENCE CHARACTERISTICS:
LENGTH: 20 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-47

Query Match 100.0%; Score 86; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 2.2e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRWPMPWRRK 12
DB 1 ILRWPMPWRRK 12

RESULT 3
US-08-915-314-46
Sequence 46, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP

STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 46:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-46

Query Match 100.0%; Score 86; DB 4; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.3e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRWPMPWRRK 12
DB 1 ILRWPMPWRRK 12

RESULT 4
US-08-915-314-48
Sequence 48, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:

TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 48:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-48

Query Match 100.0%; Score 86; DB 4; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.3e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILRPMWPMWRK 12
DB 1 ILRPMWPMWRK 12

RESULT 5
US-08-915-314-40

Sequence 40, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED AND BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 40:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-40

Query Match 96.5%; Score 83; DB 4; Length 12;
Best Local Similarity 91.7%; Pred. No. 3.5e-06;
Matches 11; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILRPMWPMWRK 12
DB 1 ILRPMWPMWRK 12

RESULT 6
US-08-915-314-76

Sequence 76, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED AND BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 76:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-76

Query Match 95.3%; Score 82; DB 4; Length 12;
Best Local Similarity 100.0%; Pred. No. 4.8e-06;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 LRPMPWPMWRK 12
DB 2 LRPMPWPMWRK 12

RESULT 7
US-08-915-314-77

Sequence 77, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED AND BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington

COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 77:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-77

Query Match 94.2%; Score 81; DB 4; Length 12;
Best Local Similarity 91.7%; Pred. No. 6.5e-06;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ILRMPWMPWRRK 12
| | | | | | | | | | | | | |
Db 1 IARMPWMPWRRK 12

RESULT 8
US-08-915-314-87
Sequence 87, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfle, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 87:

SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-87

Query Match 94.2%; Score 81; DB 4; Length 12;
Best Local Similarity 100.0%; Pred. No. 6.5e-06;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ILRMPWMPWRRK 11
| | | | | | | | | | | | | |
Db 1 ILRMPWMPWRR 11

RESULT 9
US-08-915-314-78
Sequence 78, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfle, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 78:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-78

Query Match 93.0%; Score 80; DB 4; Length 12;
Best Local Similarity 91.7%; Pred. No. 9e-06;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ILRMPWMPWRRK 12
| | | | | | | | | | | | | |
Db 1 ILRMPWMPWRRK 12

RESULT 10
US-08-915-314-85

Sequence 85, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: NO. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 85:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-85
Query Match 93.0%; Score 80; DB 4; Length 12;
Best Local Similarity 91.7%; Pred. No. 9e-06;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 ILRWPMPWRRK 12
Db 1 ILRWPMPWRRK 12
RESULT 11
US-08-915-314-86
Sequence 86, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: NO. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 86:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-86
Query Match 93.0%; Score 80; DB 4; Length 12;
Best Local Similarity 91.7%; Pred. No. 9e-06;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 ILRWPMPWRRK 12
Db 1 ILRWPMPWRRK 12
RESULT 12
US-08-915-314-80
Sequence 80, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: NO. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 80:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid

STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-80

Query Match 90.7%; Score 78; DB 4; Length 12;
Best Local Similarity 91.7%; Pred. No. 1.7e-05;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ILRPMWPMRRK 12
|||||

DB 1 ILRPMWPMRRK 12

RESULT 13
US-08-915-314-83
Sequence 83, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfle, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 83:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-83

Query Match 90.7%; Score 78; DB 4; Length 12;
Best Local Similarity 91.7%; Pred. No. 1.7e-05;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ILRPMWPMRRK 12
|||||

DB 1 ILRPMWPMRRK 12

RESULT 14
US-08-915-314-38
Sequence 38, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:

APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfle, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 38:
SEQUENCE CHARACTERISTICS:
LENGTH: 13 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-38

Query Match 90.7%; Score 78; DB 4; Length 13;
Best Local Similarity 100.0%; Pred. No. 1.9e-05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 RMPWPMRRK 12
|||||

DB 4 RMPWPMRRK 13

RESULT 15
US-08-915-314-69
Sequence 69, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfle, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/915,314
 FILING DATE: 20-AUG-1997
 CLASSIFICATION: 424
 ATTORNEY/AGENT INFORMATION:
 NAME: NO. 6180604tenburg Ph.D., Carol
 REGISTRATION NUMBER: 39,317
 REFERENCE/DOCKET NUMBER: 660081.405
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (206) 622-4900
 TELEFAX: (206) 682-6031
 INFORMATION FOR SEQ ID NO: 1:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 12 amino acids
 TYPE: amino acid
 STRANDEDNESS:
 TOPOLOGY: linear
 US-08-915-314-69

Query Match 89.5%; Score 77; DB 4; Length 12;
 Best Local Similarity 83.3%; Pred. No. 2.3e-05;
 Matches 10; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 ILRPWMPWRRK 12
 | : |||||
 Db 1 IKKPPWMPWRRK 12

Search completed: January 4, 2002, 08:40:57
 Job time: 140 sec

THIS PAGE BLANK (USPTO)


```
seq_documentation_block:
ID   AAA27298 standard; DNA; 114 BP.
XX
AC   AAA27298;
XX
DT   20-SEP-2000 (first entry)
XX
DE   Oligonucleotide used for synthesis of MBI 11B7 first cassette.
XX
KW   MBI-11; indolicidin; bovine; ss.
XX
OS   Synthetic.
XX
PN   WO200031279-A2.
XX
PD   02-JUN-2000.
XX
PE   19-NOV-1999; 99WO-CA01107.
XX
PR   20-NOV-1998; 98US-0109218.
XX
PA   (MICR-) MICROLOGIX BIOTECH INC.
XX
PI   Burian J, Bartfeld D;
XX
DR   WPI; 2000-400086/34.
XX
PT   Multi-domain fusion protein expression cassette used for high yield
PS   stable production of foreign peptide gene products -
XX
Example 5; Page 40; 73pp; English.
XX
A novel method allows the efficient production of cationic peptides in
recombinant host cells. The method involves construction of a
multi-domain fusion protein expression cassette comprising a promoter and
a nucleic acid molecule expressed as an insoluble protein. The inclusion
of anionic peptide sequences in the linker sequences neutralises the
positive charge of the cationic peptide so that the charge of the
fusion protein is controlled. This cassette allows high yield, stable
production of the cationic peptide. Cationic peptides such as
bovine indolicidin may be used as antimicrobial agents. The present
sequence is an oligonucleotide that was used in the expression
cassette. MBI-11B7 is a cationic peptide derived from modifications
of indolicidin.
XX
SQ   Sequence 114 BP; 20 A; 34 C; 32 G; 28 T; 0 other;
XX

alignment_scores:
Quality: 86.00 Length: 12
Ratio: 7.167 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-36 x AAA27298
..
Align seg 1/1 to: AAA27298 from: 1 to: 114
1 11leuAgtTrrProTrrPrrParGArgLys 12
|||||
47 ATTCTGCGTTGGCGGTGGCGGTGGCGCAAA 82

seq_name: /SID8/gcgdata/geneseq/MA2000.DAT:AAA27294
seq_documentation_block:
ID   AAA27294 standard; DNA; 151 BP.
XX
AC   AAA27294;
XX
DT   20-SEP-2000 (first entry) ;
XX
```

```
XX
DE   Oligonucleotide used for synthesis of MBI 2X11B7 last cassette.
XX
KW   Oligonucleotide; cellulose binding domain; CBD; cationic peptide;
XX   MBI-11; indolicidin; bovine; ss.
XX
OS   Synthetic.
XX
PN   WO200031279-A2.
XX
PD   02-JUN-2000.
XX
PE   19-NOV-1999; 99WO-CA01107.
XX
PR   20-NOV-1998; 98US-0109218.
XX
PA   (MICR-) MICROLOGIX BIOTECH INC.
XX
PI   Burian J, Bartfeld D;
XX
DR   WPI; 2000-400086/34.
XX
PT   Multi-domain fusion protein expression cassette used for high yield
PS   stable production of foreign peptide gene products -
XX
Example 5; Page 38; 73pp; English.
XX
A novel method allows the efficient production of cationic peptides in
recombinant host cells. The method involves construction of a
multi-domain fusion protein expression cassette comprising a promoter and
a nucleic acid molecule expressed as an insoluble protein. The inclusion
of anionic peptide sequences in the linker sequences neutralises the
positive charge of the cationic peptide so that the charge of the
fusion protein is controlled. This cassette allows high yield, stable
production of the cationic peptide. Cationic peptides such as
bovine indolicidin may be used as antimicrobial agents. The present
sequence is an oligonucleotide that was used in the expression
cassette. MBI-11B7 is a cationic peptide derived from modifications
of indolicidin.
XX
SQ   Sequence 151 BP; 22 A; 44 C; 49 G; 36 T; 0 other;
XX

alignment_scores:
Quality: 86.00 Length: 12
Ratio: 7.167 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-36 x AAA27294
..
Align seg 1/1 to: AAA27294 from: 1 to: 151
1 11leuAgtTrrProTrrPrrParGArgLys 12
|||||
41 ATTCTGCGTTGGCGGTGGCGGTGGCGCAAA 76

seq_name: /SID8/gcgdata/geneseq/MA1998.DAT:AAV60908
seq_documentation_block:
ID   AAV60908 standard; DNA; 88 BP.
XX
AC   AAV60908;
XX
DT   11-JAN-1999 (first entry)
XX
DE   DNA fragment encoding MB11.
XX
KW   MB128; cationic peptide; plasmid pK11; small cryptic plasmid;
XX   replication; RepA; vector; RAMP; human; MB11; ss.
XX
OS   Synthetic.
```



```

OS Homo sapiens.
PN MO9841636-A2.
PD 24-SEP-1998.
PF 16-MAR-1998; 98MO-CA00Q14.
PR 14-MAR-1997; 97US-00407222.
PS (BURI1) BURIAN J.
PA (KAYW/) KAY W W.
PI Burian J, Kay WW;
DR WPI: 1998-531571/45.
XX
XX Increasing plasmid copy number in a cell with the repA gene product
PT - and an small cryptic plasmid ori sequence, useful for high level
XX expression of e.g. cytokines, antigens or therapeutic proteins
XX
XX Example 16; Page 57; 82pp; English.
XX
XX This oligonucleotide was used as a template in a PCR reaction (see
CC also AAV60909-10) to generate a DNA fragment encoding the cationic
CC peptide MB11 (see AAW71690). The PCR product was cloned into
CC the universal vector pR2h-B1, which contains the R21 replication
CC leader of RepA (see AAW71686) and 2 tandem copies of the preprio
CC region (Hpro) of human defensin. The vector provides expression
CC of R21-Hpro-MB11 fusion in host cells. The invention provides a
CC controlled replication plasmid vectors (RAMP vectors) comprising a
CC replication origin of a small cryptic plasmid such as pK11 (see
CC AAV5892) and a gene encoding RepA (see AAW71686). The vectors can
CC reach very high levels of plasmid replication, but are not lethal
CC to the host cell, and can be used to direct the high level
CC expression of e.g. cytokines, antigens and therapeutic proteins.
XX
XX Sequence 88 BP; 20 A; 18 C; 25 G; 25 T; 0 other.
XX
XX
XX alignment_scores:
XX Quality: 75.00 Length: 10
XX Ratio: 7.500 Gaps: 0
XX Percent Similarity: 100.000 Percent Identity: 90.000
XX
XX alignment_block:
XX US-09-444-281-36 x AAV60908 ..
XX
XX Align seg 1/1 to: AAV60908 from: 1 to: 88
XX
XX 3 ArgTrpProTTPTrpProTTPArgAArgLys 12
XX ::::::::::::::::::::::::::::::::::::
XX 34 AATATGCGCGTGTGCGCGTGGCGTGTAA 63
XX
XX seq_name: /SIDS8/gcgdata/geneseq/geneseqn/NA2000.DAT:AAA27291
XX
XX seq_documentation_block:
XX ID AAA27291 standard; DNA; 114 BP.
XX
XX AAA27291;
XX
XX 20-SEP-2000 (first entry)
XX
XX Oligonucleotide used for synthesis of MB1-11 fragment.
DE Oligonucleotide; cellulose binding domain; CBD; cationic peptide;
XX MB1-11; indolicidin; bovine; ss.
XX
XX Synthetic.
XX
XX WO200031279-A2.
XX
XX 02-JUN-2000.
XX

```

```

XX PF 19-NOV-1999;      99WO-CA01107.
XX XX 20-NOV-1998;     98US-0109218.
XX PR
XX XX
XX PA (MCCR-) MICROLOGIX BIOTECH INC.
XX PI Burian J, Bartfield D;
XX DR WPI; 2000-400086/34.
XX XX
PT PT Multi-domain fusion protein expression cassette used for high yield
PS stable production of foreign peptide gene products -
PS Example 4; Page 37; 73pp; English.
XX XX
CC CC A novel method allows the efficient production of cationic peptides in
CC recombinant host cells. The method involves construction of a
CC multi-domain fusion protein expression cassette comprising a promoter an
CC a nucleic acid molecule expressed as an insoluble protein. The inclusion
CC of anionic peptide sequences in the linker sequences neutralises the
CC positive charge of the cationic peptide so that the charge of the
CC fusion protein is controlled. This cassette allows high yield, stable
CC production of the cationic peptide. Cationic peptides such as
CC bovine indolicidin may be used as antimicrobial agents. The present
CC sequence is an oligonucleotide that was used to synthesise a
CC MBI-11 fragment. MBI-11 is a cationic peptide derived from modifications
CC of indolicidin.
XX CC
SQ Sequence 114 BP; 25 A; 26 C; 30 G; 33 T; 0 other;

alignment_scores:
    Quality:   75.00          Length:       10
    Ratio:     7.500         Gaps:        0
    Percent Similarity: 100.000    Percent Identity: 90.000

alignment_block:
US-09-444-281-36 x AAA27291 ..

Align seg 1/1 to: AAA27291 from: 1 to: 114

      3 ArgTTPROTTRPTrPPROTPARgArgLys 12
      ::::::::::::::::::::
      50 AAATGCGCTGGTGCGCCGTGCCTGTAA 79

seq_name= /SIDS8/gcgdata/geneseq/geneseqn/NA1999.DAT:AAV83788

seq_documentation_block:
ID ID AAV83788 standard; DNA; 39 BP.
XX AC AAV83788;
XX XX
DT DT 19-MAR-1999 (first entry)
XX DE
DE Antimicrobial peptide Indolicidin encoding DNA.
XX KW Antimicrobial; fusion; acidic peptide; recombinant; microorganism;
KW guanamerin; basic peptide; indolicidin; ss.
XX OS Synthetic.
OS Bos sp.
XX FH
FH Key Location/Qualifiers
FT CDS 1..39
FT /*lag= a
FT ./note= "the start and stop codons are not indicated"
XX PN WO9854336-A1.
XX PD
PD 03-DEC-1998.
XX XX
XX 28-MAY-1998; 98MO-KR00132.
```

```
XX 09-APR-1998; 98KR-0013372.
PR 28-MAY-1997; 97KR-0021312.
XX
XX (KOAD ) KOREA ADV INST SCI & TECHNOLOGY.
PA (SAMY-) SAMYANG GENEX CORP.
XX
XX Hong S, Kang MH, Kim JH, Kim S, Lee H, Lee JH;
XX WPI; 1999-059844/05.
DR P-PSDB; AAW87609.
XX
PT New method for mass production of antimicrobial peptides - by
PT constructing fusion genes comprising acidic and antimicrobial
PT peptide genes and transforming host with vector containing these
XX
XX Example 6; Page 18; 52pp; English.
XX
XX The invention relates to mass production of antimicrobial peptides. The
XX method comprises constructing a fusion gene containing a first gene
XX encoding a negatively charged acidic peptide having at least two cysteine
XX residues, and a second gene encoding a positively charged basic
XX antimicrobial peptide. A host microorganism is transformed with a vector
XX containing the fusion gene and then cultured. The expressed antimicrobial
XX peptide is then recovered. The method is used to mass produce
XX antimicrobial peptides in recombinant microorganisms. The inhibitory
XX effect of the expressed antimicrobial peptide upon the growth of the host
XX microorganism is considerably reduced by fusing it to the acidic peptide.
XX Therefore, the use of the fusion gene provides an economic, recombinant
XX alternative of mass producing antimicrobial peptides, which overcomes the
XX disadvantages of low-productivity and poor economy, previously
XX encountered by recombinant and chemical methods. The present sequence
XX represents the DNA encoding an antimicrobial peptide indolicidin. This
XX can be used along with the acidic peptide Guamerin gene in the
XX construction of the fusion gene.
XX
SQ Sequence 39 BP; 4 A; 10 C; 16 G; 9 T; 0 other;

alignment_scores:
      Quality: 70.00      Length: 9
      Ratio: 7.778      Gaps: 0
Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:
US-09-444-281-36 x AAV83788 ..

Align seq 1/1 to: AAV83788 from: 1 to: 39
      3 ArgTrpProTrrPrpGTPARG 11
      ::::::::::::::::::::11
      13 AATGCGCGTGTGGCGCGTGTGT 39

seq_name: /SID8/gcgdata/geneseq/NA2000.DAT:AAZ29389

seq_documentation_block:
ID AAZ29389 standard; DNA; 47 BP.
XX
XX AAZ29389;
XX
XX 29-FEB-2000 (first entry)
XX
XX PCR primer-15 for synthesis of antimicrobial peptide indolicidin gene.
XX
XX PCR primer: anti-microbial peptide; indolicidin gene; DNA construct;
XX glutamine pyrophosphoribosyl pyrophosphatase gene;
XX purf gene; fusion peptide; mass production; pharmaceutical industry;
XX food industry; ss.
XX
XX Synthetic.
XX
XX W0964611-A1.
XX
```

```
PD 16-DEC-1999.
XX
XX 08-JUN-1999; 99WO-KR00282.
XX
XX 09-JUN-1998; 98KR-0022117.
PR 14-MAY-1999; 99KR-0017920.
XX
XX (SAMY-) SAMYANG GENEX CORP.
XX
XX Kim JH, Kang MH, Lee J, Park SH, Lee JW, Hong SS, Lee H;
XX WPI; 2000-097542/08.
DR
XX
XX New DNA constructs useful for mass production of antimicrobial peptides
XX in microorganism hosts -
XX
XX Example 1; Page 13; 67pp; English.
XX
XX The present sequence is a chemically synthesised PCR primer which was
XX used to synthesise a gene encoding antimicrobial peptide indolicidin.
XX The antimicrobial peptide gene is used in a DNA construct that comprises
XX entire, partial or a derivative of purf gene (glutamine
XX pyrophosphoribosyl pyrophosphatase gene). The DNA
XX construct allows mass production of the antimicrobial peptide in
XX microbial hosts without killing the host cells. The antimicrobial
XX peptides are useful commercially in the pharmaceutical and
XX food industries.
XX
SQ Sequence 47 BP; 6 A; 11 C; 19 G; 11 T; 0 other;

alignment_scores:
      Quality: 70.00      Length: 9
      Ratio: 7.778      Gaps: 0
Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:
US-09-444-281-36 x AAZ29389 ..

Align seq 1/1 to: AAZ29389 from: 1 to: 47
      3 ArgTrpProTrrPrpGTPARG 11
      ::::::::::::::::::::11
      17 AATGCGCGTGTGGCGCGTGTGT 43

seq_name: /SID8/gcgdata/geneseq/NA2000.DAT:AAZ29390

seq_documentation_block:
ID AAZ29390 standard; DNA; 47 BP.
XX
XX AAZ29390;
XX
XX 29-FEB-2000 (first entry)
XX
XX PCR primer-16 for synthesis of antimicrobial peptide indolicidin gene.
XX
XX PCR primer: anti-microbial peptide; indolicidin gene; DNA construct;
XX glutamine pyrophosphoribosyl pyrophosphatase gene;
XX purf gene; fusion peptide; mass production; pharmaceutical industry;
XX food industry; ss.
XX
XX Synthetic.
XX
XX W0964611-A1.
XX
XX 16-DEC-1999.
XX
XX 08-JUN-1999; 99WO-KR00282.
XX
XX 09-JUN-1998; 98KR-0022117.
PR 14-MAY-1999; 99KR-0017920.
XX
XX (SAMY-) SAMYANG GENEX CORP.
XX
```

PI Kim JH, Kang MH, Lee J, Park SH, Lee JW, Hong SS, Lee H;

PT Inhibiting growth of sulphate-reducing bacteria using other bacteria
PT particularly for protection of metals and concrete -
XX

PS Example 4; Fig 1; 84pp; English.
 CC This sequence represents an oligonucleotide for cloning the non-amidated
 CC indolicidin peptide coding sequence. The invention relates to a method
 CC for inhibiting growth of sulphate-reducing bacteria (A) on a material (B)
 CC sensitive to corrosion or degradation, by applying to (B) a bacterium (C)
 CC that secretes a compound (I) able to inhibit growth of (A). The method is
 CC used to protect metal, concrete or cement against corrosion and
 CC degradation, but (B) can also be used to protect dental implants, (B) is
 CC present in an open or closed system (e.g. water cooling tower, liquid
 CC storage container, fuel tank, sewer or drainage system etc.) or part of a
 CC bridge or other structure. The method is more effective and less
 CC expensive than known methods for inhibiting (A), and reduces the amount
 CC of toxic chemicals released. Conventional biofilms of aerobic organisms
 CC tend to encourage growth of (A), and addition of (C) to the biofilm
 CC prevents this. A single application of (C) lasts for a long time, and (I)
 CC are produced exactly where they are required and inhibit (A) without
 CC significant impact on other organisms (this effect includes reducing
 CC resistance of (A) to conventional biocides, which may then be used in
 CC reduced amounts). If local damage to the biofilm occurs, the underlying
 CC material is still protected by diffusion of (I) from neighbouring areas.
 CC
 SQ Sequence 69 BP; 14 A; 18 C; 20 G; 17 T; 0 other;

alignment_scores:
 Quality: 70.00 Length: 9
 Ratio: 7.778 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:
 US-09-444-281-36 x AAZ40246 ..

Align seg 1/1 to: AAZ40246 from: 1 to: 69

```

3 ArgTrpProTyrProTyrPArgArg 11
:::|||||
28 AAATGGCCTTGTCGCTTGCGCCGC 54

```

seq_name: /SIDSB/gcgdata/geneseq/geneseqn/NA2000.DAT:AAZ49764

seq_documentation_block:
 ID AAZ49764 standard; DNA; 211 BP.

AC AAZ49764;

DT 18-APR-2000 (first entry)

DE Poly-(Indol (1-13)-Met-Ala-AIy-Ile-Ala-Met)3 DNA.

XX Crosslinked indolicidin analog; X-indolicidin; poly-indol 1-13;
 KW stability; bovine neutrophil; antimicrobial; antibacterial; fungicide;
 KW protozoacide; virucide; anti-HIV; human immunodeficiency virus-1;
 KW HIV-1; gram positive bacteria; gram negative; Staphylococcus aureus;
 KW Escherichia coli; Salmonella typhimurium; yeast; fungi; protozoa;
 KW Candida albicans; Cryptococcus neoformans; Giardia; Acanthamoeba;
 KW hexapeptide spacer; ds.
 XX

OS Synthetic.

OS Bos sp.

XX Key Location/Qualifiers
 XX CDS 8..199

FT /tag= a
 FT /product= "Poly-(Indol(1-13)-Met-Ala-Arg-Ile-Ala-Met)3"
 FT /note= "encodes three copies of Indol 1-13, each
 FT separated by Met-Ala-Arg-Ile-Ala-Met spacer sequence"

FT primer_bind
 FT 1..21
 FT /tag= b

FT primer_bind
 FT complement (191..211)
 FT /tag= c

FT misc_feature
 FT 68..71
 FT /tag= d

FT

FT /note= "corresponds to overlap in oligonucleotides
 FT used for ligation"
 FT 148..151
 FT misc_feature
 FT /tag= e
 FT /note= "corresponds to overlap in oligonucleotides
 FT used for ligation"
 PN W09965510-A1.
 PD 23-DEC-1999.
 XX
 XX 20-MAY-1999; 99MO-US11165.
 XX
 XX 18-JUN-1998; 98US-0099631.
 XX
 XX (REGC) UNIV CALIFORNIA.
 XX
 XX Selsled ME, Osapay K;
 PI
 XX WPI; 2000-147133/13.
 DR P-PSDB; AAY44668.
 XX

PS Crosslinked indolicidin analogs with antimicrobial activity against
 PT bacteria, yeast, fungi, protozoa and viruses
 XX

XX Example 1C; Fig 1; 53pp; English.
 CC The patent discloses crosslinked analogs of indolicidin (indol 1-13)
 CC which is a naturally occurring peptide isolated from bovine neutrophils
 CC and has antimicrobial activity. The crosslinked indolicidin
 CC (X-indolicidin) analogs are stable and have antimicrobial activity
 CC against gram positive and negative bacteria (e.g. Staphylococcus aureus,
 CC Escherichia coli and Salmonella typhimurium), yeasts and fungi (e.g.
 CC Candida albicans, Cryptococcus neoformans), protozoa (e.g. Giardia
 CC species and Acanthamoeba species), and viruses (e.g. HIV-1).
 CC They can be used for reducing or inhibiting the growth or survival of
 CC microorganisms in an environment e.g. a food or food product, a
 CC solution, an inanimate object comprising a surface, or a mammal.
 CC The present sequence is a DNA encoding a protein comprising three
 CC copies of indol 1-13 each separated by a hexapeptide spacer sequence.
 CC The sequence was used to produce a recombinant construct for the
 CC expression of indol-homoserine (Hse) analog. The ability of
 CC indol-Hse analog to maintain antimicrobial activity provides a means to
 CC produce X-indolicidin analog precursors in sufficient quantities.
 CC
 SQ Sequence 211 BP; 36 A; 50 C; 74 G; 51 T; 0 other;

alignment_scores:
 Quality: 70.00 Length: 9
 Ratio: 7.778 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:
 US-09-444-281-36 x AAZ49764 ..

Align seg 1/1 to: AAZ49764 from: 1 to: 211

```

3 ArgTrpProTyrProTyrPArgArg 11
:::|||||
38 AAATGGCCTTGTCGCTTGCGCTGCT 64

```

seq_name: /SIDSB/gcgdata/geneseq/geneseqn/NA2000.DAT:AAZ45123

seq_documentation_block:
 ID AAZ45123 standard; DNA; 211 BP.

AC AAZ45123;

DT 28-FEB-2000 (first entry)

DE Indolicidin fusion peptide nucleotide sequence.

XX

Indolicidin analogue; antimicrobial activity; helminth; bacteria; virus;
treatment; inhibit growth; micro-organism; contact lens solution;
transgenic plant; surgical instrument; yeast; fungi; protozoa; ss.
XX
OS Synthetic.
XX
PN M09958141-A1.
XX
PD 18-NOV-1999.
XX
PE 05-MAY-1999; 99WO-US09942.
XX
PR 12-MAY-1998; 98US-0076227.
XX
PA (REGC) UNIV CALIFORNIA.
XX
PI Seasted ME;
XX
DR WPI; 2000-053028/04.
XX
P-PSDB; AAY57142.
XX
PT New indolicidin analogues, active against bacteria, yeast, fungi,
XX
PS protozoa and virus, used for, e.g. treating infections.
XX
PS Disclosure; Fig 6; 62pp; English.
XX
CC This is the nucleotide sequence of an example of a fusion protein which
CC consists of an indolicidin analogue linked to another peptide.
CC Peptides AAY57109-Y57138 and AAY57143-Y57144 are new indolicidin
CC analogues, which have a homoserine residue and/or a truncated amino
CC terminal region. The analogues have the following amino acid sequence:
CC Xaa1-Xaa2-Xaa3-Xaa4-Xaa5-Xaa6-Pro-Xaa6-Xaa6-Pro-Xaa6-Xaa7-Xaa8
CC where:
CC Xaa1 = Ile, Leu, Val, Ala, Gly or absent;
CC Xaa2 = Ile, Leu, Val, Ala, Gly or absent;
CC Xaa3 = Pro or absent;
CC Xaa4 = Trp, Phe or absent;
CC Xaa5 = Arg, Lys or absent;
CC Xaa6 = Trp or Phe;
CC Xaa7 = Arg, Lys or absent;
CC Xaa8 = homoserine (Hse), Met, Met-Xaa9-Met or absent, and
CC Xaa9 = at least one amino acid;
CC provided that if Xaa1 is present, Xaa8 = Hse, Met or Met-Xaa9-Met;
CC and further provided that: if Xaa2 is absent, Xaa1 is absent; if Xaa3 is
CC absent, Xaa1 and Xaa2 are absent; if Xaa4 is absent, Xaa1 and Xaa2
CC are absent; and if Xaa5 is absent, Xaa1, Xaa2, Xaa3 and Xaa4 are absent.
CC The indolicidin analogues can be used to create a fusion polypeptide
CC consisting of the analogue linked to a peptide. The indolicidin
CC analogues have antimicrobial activity against gram positive bacteria,
CC gram negative bacteria, yeast, fungus, protozoa and viruses (e.g. HIV-1).
CC They are also active against helminths. The analogues can be used for
CC reducing or inhibiting growth or survival of a microorganism. They can be
CC used for treating infections. They can also be included in a liquid such
CC as water or an aqueous solution, e.g. contact lens solution. The
CC analogues have potential uses in food products, and in objects such as
CC the surface of an instrument used to prepare food or to perform surgery.
CC Transgenic plants or animals useful in the food industry can be produced
CC by introducing a nucleic acid molecule encoding an indolicidin analogue
CC into the germ-line cells of such organisms.
XX
XX Sequence 211 BP; 36 A; 50 C; 74 G; 51 T; 0 other;

alignment_scores:
Quality: 70.00 Length: 9
Ratio: 7.778 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:
US-09-444-281-36 x AAZ45123

Align seg 1/1 to: AAZ45123 from: 1 to: 211

3 ArgTrpProTTrpProTrpArgArg 11
:::|||||
38 AATGGCGCTGCTGCGCTGCGCTGCT 64

seq_name: /SID58/gcdata/geneseq/geneseq/NA1999.DAT:AAZ20646

seq_documentation_block:

ID AAZ20646 standard; RNA; 6446 BP.

AAZ20646;

26-NOV-1999 (first entry)

TMV-based virus TMV861 coat protein read-through RNA sequence.

TMV-based virus; tobacco mosaic virus; protein isolation; green juice;

virus isolation; fusion protein identification; ss.

Tobacco mosaic virus.

W09946288-A2.

16-SEP-1999.

09-MAR-1999; 99WO-US05056.

10-MAR-1998; 98US-0037751.

(BIOS-) BIOSOURCE TECHNOLOGIES INC.

Gargier SJ, Holtz RB, McCulloch MJ, Turpen TH;

WPI; 1999-561660/47.

Obtaining protein, viruses and fusion proteins from plants, using
non-denaturing conditions

Disclosure; Page 55-58; 58pp; English.

This sequence represents a tobacco mosaic virus (TMV) based virus
sequence identified using the method of the invention. The method is for
obtaining a soluble protein or peptide of interest from a plant,
comprising homogenising the plant to produce green juice, adjusting the
pH to less than or equal to 5.2, and heating the juice to a minimum of
45 degrees C. The juice is then centrifuged to produce a supernatant, and
the protein or peptide is purified from the supernatant. The method can
also be used for obtaining viruses and fusion proteins. The method is
especially useful for obtaining IL-1 to IL-10, Epo, G-CSF, GM-CSF,
hP-CSF, M-CSF, Factor VIII, Factor IX, tPA, receptors, receptor
antagonists, antibodies, single-chain antibodies, enzymes,
neuropolypeptides, insulin, antigens, vaccines, peptide hormones,
calcitonin, and human growth hormone, or an antimicrobial peptide or
protein from protegrins, magainins, cecropins, melittins, indolicidins,
defensins, beta-defensins, cryptidins, clavinins, plant defensins,
nicin and bactericins, all produced by recombinant means. The new method
is more efficient than the prior art for isolating viruses, protein, and
peptides. The method is large-scale, and non-denaturing and
solvent-limited. Prior art methods do not isolate recombinant proteins,
and do not allow fraction 2 proteins to be ultrafiltrated.

Sequence 6446 BP; 1873 A; 1234 C; 1563 G; 1776 U; 0 other;

alignment_scores:
Quality: 70.00 Length: 9
Ratio: 7.778 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:
US-09-444-281-36 x AAZ20646

Align seg 1/1 to: AAZ20646 from: 1 to: 6446

3 ArgTrpProTrrPProTrrPArg 11
:::|||||
6213 AAGUGGCCUUGGCGCAUGGCGCGA 6239

seq_name: /SID58/gcgdata/geneseq/geneseq/NA2001.DAT:AAF82334

seq_documentation_block:

ID AAF82334 standard; RNA: 6446 BP.

AAF82334:

22-JUN-2001 (first entry)

Tobacco mosaic virus-based coat protein read-through virus TMW861.

TMW: tobacco mosaic virus; TMW861: virus isolation;

KW non-native protein purification; ribulose 1,5-diphosphate carboxylase;

KM Rubisco; coat protein read-through; ss.

Tobacco mosaic virus.

WO200119969-A1.

22-MAR-2001.

19-MAY-2000; 2000WO-US13680.

16-SEP-1999; 99US-0397090.

(LARG-) LARGE SCALE BIOLOGY CORP.

Garger SJ, Holtz BR, McCulloch MJ, Turpen TH;

WPI; 2001-328016/34.

Minimizing presence of ribulose 1,5-diphosphate carboxylase to obtain

plant product for isolating bioactive species. Involves cutting plant

material from plant in cutting period when quantity of Rubisco is at

minimum

Disclosure; Page 72-74; 81pp; English.

The present sequence is a tobacco mosaic virus (TMV)-based virus

which was used to infect field-grown tobacco. The virus was then

isolated from the tobacco plants by a novel process for isolating and

purifying viruses, soluble proteins and peptides from plant sources. In

order to isolate the bioactive species from the undesirable

photosynthetic protein ribulose 1,5-diphosphate carboxylase (Rubisco),

the plant material is cut in a period of the light/dark cycle when the

quantity of Rubisco in the plant is at a minimum. The method is useful

for obtaining a virus of interest. It is also useful for obtaining

soluble recombinant or non-native proteins, such as active mammalian

proteins, enzymes, vaccines, antibodies and peptides, from transgenic

plants.

Sequence 6446 BP; 1873 A; 1234 C; 1563 G; 1776 U; 0 other;

alignment_scores:

Quality: 70.00 Length: 9.

Ratio: 7.778 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:

US-09-444-281-36 x AAF82334

Align seg 1/1 to: AAF82334 from: 1 to: 6446

3 ArgTrpProTrrPProTrrPArg 11

:::|||||

6213 AAGUGGCCUUGGCGCAUGGCGCGA 6239

seq_name: /SID58/gcgdata/geneseq/geneseq/NA2000.DAT:AAA28519

seq_documentation_block:

ID AAA28519 standard; DNA: 207 BP.

AAA28519:

29-AUG-2000 (first entry)

PCRIL DNA coding sequence.

Magainin; antimicrobial; transgenic plant; protease degradation; Rev4;

indolicidin; protein production; reverse peptide; ss.

Synthetic.

WO200026344-A1.

11-MAY-2000.

29-OCT-1999; 99WO-US25561.

30-OCT-1998; 98US-0106373.

02-NOV-1998; 98US-0106373.

(INTE-) INTERLINK BIOTECHNOLOGIES LLC.

(KENT) UNIV KENTUCKY RES FOUND.

Everett NP, LI Q, Lawrence C, Davies MH;

WPI; 2000-365597/31.

P-PSDB; AAY92840.

Polypeptides for reducing proteolytic degradation of proteins

administered to, or produced by a plant comprise indolicidin or its

functional equivalents

Example 17; Page 35; 50pp; English.

Indolicidin is a potent antimicrobial tridecapeptide, originally

purified from cytoplasmic granules of bovine neutrophils. Reverse

peptide, Rev4 of indolicidin (see AAY92794) was found to have increased

stability against plant protease degradation. Expression of antimicrobial

peptides in transgenic plants suffers a major limitation in that the

foreign peptides are susceptible to rapid degradation by proteases. The

invention concerns reducing the extent of protease degradation of a

protein applied to, or produced by a plant by administering indolicidin,

Rev4 or a functional equivalent to the plant. Transgenic plants

expressing indolicidin and Rev4 are useful for production of the

antimicrobial peptides. Compositions containing indolicidin and Rev4 are

also useful for production of agronomically important proteins in

plants.

Sequence 207 BP; 49 A; 50 C; 36 G; 72 T; 0 other;

alignment_scores:

Quality: 67.00 Length: 10

Ratio: 7.444 Gaps: 0

Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:

US-09-444-281-36 x AAA28519

Align seg 1/1 to: AAA28519 from: 1 to: 207

1 IleleuArgTrpProTrrPProTrrPArg 10

||| |||

163 ATTAGAGATGCGCTTGCTGCTTGGA 192

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:40:26 : Search time 53.46 Seconds
(without alignments)
18.013 Million cell updates/sec

Title: US-09-444-281-35
Perfect score: 91
Sequence: 1 ILKMPMPWRRK 13

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 522463 seqs, 74073290 residues
Total number of hits satisfying chosen parameters: 522463

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

1: A_Geneseq_1101.*
2: /SIDSB/gcgdata/geneseq/geneseq/AA1980.DAT.*
3: /SIDSB/gcgdata/geneseq/geneseq/AA1981.DAT.*
4: /SIDSB/gcgdata/geneseq/geneseq/AA1982.DAT.*
5: /SIDSB/gcgdata/geneseq/geneseq/AA1983.DAT.*
6: /SIDSB/gcgdata/geneseq/geneseq/AA1984.DAT.*
7: /SIDSB/gcgdata/geneseq/geneseq/AA1985.DAT.*
8: /SIDSB/gcgdata/geneseq/geneseq/AA1986.DAT.*
9: /SIDSB/gcgdata/geneseq/geneseq/AA1987.DAT.*
10: /SIDSB/gcgdata/geneseq/geneseq/AA1988.DAT.*
11: /SIDSB/gcgdata/geneseq/geneseq/AA1989.DAT.*
12: /SIDSB/gcgdata/geneseq/geneseq/AA1990.DAT.*
13: /SIDSB/gcgdata/geneseq/geneseq/AA1991.DAT.*
14: /SIDSB/gcgdata/geneseq/geneseq/AA1992.DAT.*
15: /SIDSB/gcgdata/geneseq/geneseq/AA1993.DAT.*
16: /SIDSB/gcgdata/geneseq/geneseq/AA1994.DAT.*
17: /SIDSB/gcgdata/geneseq/geneseq/AA1995.DAT.*
18: /SIDSB/gcgdata/geneseq/geneseq/AA1996.DAT.*
19: /SIDSB/gcgdata/geneseq/geneseq/AA1997.DAT.*
20: /SIDSB/gcgdata/geneseq/geneseq/AA1998.DAT.*
21: /SIDSB/gcgdata/geneseq/geneseq/AA1999.DAT.*
22: /SIDSB/gcgdata/geneseq/geneseq/AA2000.DAT.*
23: /SIDSB/gcgdata/geneseq/geneseq/AA2001.DAT.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	91	100.0	13	18 AAW12873	Antimicrobial catl
2	91	100.0	13	19 AAY24609	Indolicidin analog
3	91	100.0	13	19 AAW6378	Cationic peptide o
4	91	100.0	13	19 AAW71690	Cationic peptide M
5	91	100.0	13	21 AAY94495	MBI-11 peptide der
6	91	100.0	13	21 AAY92795	Indolicidin analog
7	91	100.0	13	21 AAY91773	Amino acid sequenc
8	91	100.0	13	21 AAY91774	Amino acid sequenc
9	91	100.0	13	21 AAY91818	Amino acid sequenc
10	91	100.0	13	21 AAY91819	Amino acid sequenc
11	91	100.0	13	21 AAY91820	Amino acid sequenc

12	91	100.0	14	19 AAY24583	Indolicidin analog
13	91	100.0	14	21 AAY91811	Amino acid sequenc
14	91	100.0	21	19 AAY24582	Indolicidin analog
15	91	100.0	21	21 AAY91806	Amino acid sequenc
16	87	95.6	12	21 AAY24580	Indolicidin analog
17	87	95.6	12	21 AAY91804	Amino acid sequenc
18	86	94.5	12	18 AAM12877	Antimicrobial catl
19	86	94.5	12	18 AAM12877	Antimicrobial catl
20	86	94.5	12	19 AAY24615	Indolicidin analog
21	86	94.5	13	21 AAY91833	Amino acid sequenc
22	86	94.5	13	21 AAY24572	Indolicidin analog
23	86	94.5	13	21 AAY91812	Amino acid sequenc
24	86	94.5	14	19 AAY24573	Indolicidin analog
25	86	94.5	15	18 AAY91813	Amino acid sequenc
26	86	94.5	20	19 AAM13802	Antimicrobial catl
27	86	94.5	20	21 AAY24570	Indolicidin analog
28	86	94.5	21	21 AAY91807	Amino acid sequenc
29	86	94.5	21	19 AAY24571	Indolicidin analog
30	85	93.4	12	19 AAY91808	Amino acid sequenc
31	85	93.4	12	21 AAY24586	Indolicidin analog
32	85	93.4	13	18 AAY91828	Amino acid sequenc
33	85	93.4	13	18 AAM27179	Antimicrobial catl
34	85	93.4	13	18 AAM12869	Antimicrobial catl
35	85	93.4	13	19 AAM12894	Antimicrobial catl
36	85	93.4	13	19 AAY24610	Indolicidin analog
37	85	93.4	13	21 AAY24565	Indolicidin analog
38	85	93.4	13	21 AAY91786	Amino acid sequenc
39	83	91.2	12	19 AAY91795	Amino acid sequenc
40	83	91.2	12	19 AAY24568	Indolicidin analog
41	83	91.2	12	21 AAY91789	Amino acid sequenc
42	83	91.2	13	18 AAM12892	Antimicrobial catl
43	83	91.2	13	18 AAM12893	Antimicrobial catl
44	83	91.2	13	18 AAM12896	Antimicrobial catl
45	83	91.2	13	19 AAM12897	Antimicrobial catl
			19	AA124612	Indolicidin analog

ALIGNMENTS

RESULT 1	
AAW12873	
ID AAW12873 standard; peptide; 13 AA.	
AC AAW12873:	
XX	
DT 10-DEC-1997 (first entry)	
XX	
DE Antimicrobial cationic peptide CP-11.	
XX	
KW Bacterial; viral; antitumour; food; preservative; inhibitor; growth;	
KW bacterium; yeast; endotoxaemia; sepsis; antibiotic; fungal;	
KW antiviral; candida albicans; steriliant; salmonella; yersinia;	
KW shigella.	
XX	
OS Synthetic.	
XX	
PN W09708199-A2.	
XX	
PD 06-MAR-1997.	
XX	
PF 23-AUG-1996; 96MO-IB00996.	
XX	
PR 23-AUG-1995; 95US-0002687.	
XX	
PA (UYBR-) UNIV BRITISH COLUMBIA.	
XX	
PI Falla TJ, Cough M, Hancock RW;	
XX	
DR WPI: 1997-179179/16.	
XX	
PT Cationic peptide(s) having anti-microbial activity - used for the	
PT inhibition of bacterial and viral growth, as an antitumour agent,	
PT and as a food preservative	

XX Claim 2; Page 65; 89pp; English.

XX The present sequence represents a specifically claimed novel isolated
XX cationic peptide which has antimicrobial activity. The amino acid
XX sequence of antimicrobial cationic peptides (including the present
XX sequence) is selected from: XIX1ProX2X3X2Pro(X2X2Pro)X2X3(X5)O;
XX XIX1ProX2X3X4(X5)ProX2X3X3; XIX1X3(ProTrp)uX3X2X5X2X5X2(X5)O;
XX XIX1X3X3X2X2Pro(X2X2Pro)X2(X5)m; where m = 1-5; n = 1-2; o = 2-5; r
XX = 0-8; u = 0-1; X1 = Ile, Leu, Val, Phe, Tyr, Trp or Met; X2 = Trp or
XX Phe; X3 = Arg or Lys; X4 = Trp or Lys; and X5 = Phe, Trp, Arg, Lys or
XX Pro. The peptides are preferably amidated or carboxymethylated. The
XX peptides may be used in methods for inhibiting the growth of a bacterium
XX or yeast, or for inhibiting an endotoxaemia or sepsis associated
XX disorder in a subject. The peptides have a broad activity against
XX antibiotic resistant bacteria, combined with activity against the
XX medically important fungus *Candida albicans*. In addition, the peptides
XX are useful as antileumour agents and/or antiviral agents. The peptides
XX may be used as sterilants or preservatives of materials susceptible to
XX microbial or viral contamination, e.g. in processed foods to inhibit
XX *Salmonella*, *Yersinia* and *Shigella*. The peptides are compact and tend to
XX have a unique polypyrrolone type II extended helix structure that permits
XX them to span the membrane with relatively few amino acids. The peptides
XX possess the ability to work synergistically with antibiotics, and in
XX addition, some of them possess anti-endotoxin activity.
XX N.B. The present sequence represents SEQ ID NO:1 in the claims and
XX examples of the specification, but differs slightly from the SEQ ID NO:1
XX in the sequence listing on page 51 of the specification (see AAW21719).

Sequence 13 AA;

Query Match 100.0%; Score 91; DB 18; Length 13;

Best Local Similarity 100.0%; Pred. No. 7.2e-07;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKKWPMWPMRRK 13
| | | | | | | | | | | | | | |
Db 1 ILKKWPMWPMRRK 13

RESULT 2

AAW24609
ID AAW24609 standard; peptide; 13 AA.

AC AAW24609;

DT 18-AUG-1999 (first entry)

DE Indolicidin analogue #61.

XX Indolicidin: bacterial infection; photo-oxidised solubilisier;
XX antimicrobial; antibiotic; antiarrhythmic; surface disinfectant;
XX additive; shampoo; soap; insecticide; herbicide; preservative;
XX food; technical material.

OS Synthetic.

PN WO9807745-A2.

PD 26-FEB-1998.

PF 21-AUG-1997; 97WO-0514779.

PR 13-JAN-1997; 97US-0034949.

PR 21-AUG-1996; 96US-0024754.

PA (MICR-) MICROLOGIX BIOTECH INC.

PI Erle D, Fraser JR, Krieger TJ, Taylor R, West MH;
DR WPI, 1998-169090/15.
XX

PT New indolicidin analogues with antimicrobial activity and related
PT nucleic acid - vectors, transformed cells and antibodies, also
PT conjugates with polyoxyalkylene glycol and fatty acid to reduce
PT toxicity, useful therapeutically, as disinfectants etc.

PS Example 1; Page 32; 129pp; English.

XX AAW24549 to AAW24615 represent indolicidin analogues of formulae
XX (I)-(VIII) containing up to 25 amino acids (aa): RZXXZXB (I), BXZXXZXB
XX (II), BBXZXXZXB (III), BXZXXZBBn(AA)nILBBACS (IV), BXZXXZBB(AA)nM
XX (V), LBnXZnXZnXnXnR (VI), LKXZXXZXXRRK (VII) and BBXZXXZBBB (VIII).
XX where Z = P or V; X = hydrophobic residue, preferably W; B = basic aa,
XX preferably R or K; AA = any aa; n = 0 or 1; in (II), at least 1 Z = V;
XX in (VIII) at least 2 X = P or Y. The analogues are used to treat
XX infections caused by bacteria (Gram positive or negative, or anaerobic);
XX fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
XX trematodes) or viruses. Typical of very many pathogens that can be
XX controlled are *Leishmania*, *Trypanosoma*, *Ascaris lumbricoides*, *Fasciola*
XX *hepatica*, *Listeria*, *Clostridium*, *rotavirus* and *papilloma virus*. Compounds
XX derived from the analogues may be used similarly; the compounds may
XX also be prepared from antibiotics or antiarrhythmic agents. The analogues
XX may be used therapeutically or to coat medical devices; also they are
XX useful as surface disinfectants, as additives to shampoo or soaps, as
XX insecticides or herbicides, or as preservatives for foods and technical
XX materials. The analogues are administered by injection, lavage, orally
XX or topically, generally at 0.1-50 mg/kg. These analogues have a broader
XX spectrum of activity than indolicidin and modification as compounds
XX reduces their toxicity.

Sequence 13 AA;

Query Match 100.0%; Score 91; DB 19; Length 13;

Best Local Similarity 100.0%; Pred. No. 7.2e-07;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKKWPMWPMRRK 13
| | | | | | | | | | | | | | |
Db 1 ILKKWPMWPMRRK 13

RESULT 3

AAW6378
ID AAW6378 standard; peptide; 13 AA.

AC AAW6378;

DT 12-JAN-1999 (first entry)

DE Cationic peptide of claim 15 #5.

XX Indolicidin analogue; resistance; cationic peptide; antibiotic;
XX bacterial infection; tolerance; antibacterial; microorganism;
XX bacteria; fungus; parasite; virus.

OS Synthetic.

PN WO9840401-A2.

PD 17-SEP-1998.

PF 10-MAR-1998; 98WO-CA00190.

PR 25-FEB-1998; 98US-0030619.

PR 10-MAR-1997; 97US-0040649.

PR 20-AUG-1997; 97US-0915314.

PR 26-SEP-1997; 97US-0060099.

PA (MICR-) MICROLOGIX BIOTECH INC.

PI Fraser JR, McNicol PJ, West MHP;
XX

DR WPI: 1998-520800/44.
XX New indolicidin peptide analogues, useful for, e.g. enhancing
PT activity of antibiotic or overcoming tolerance, acquired resistance
PT or inherent resistance of microorganisms
XX
PS Claim 15: Page 93; 105pp; English.
XX
CC The present sequence represents a specifically claimed cationic peptide
CC from the present invention. The present invention describes compositions
CC and methods for treating infection, especially bacterial infections. The
CC compositions and methods use cationic peptides in combination with an
CC antibiotic agent which are then administered to a patient to enhance the
CC activity of the antibiotic agent, to overcome: (a) tolerance; (b)
CC acquired resistance; and (c) inherent resistance. The combinations of
CC antibiotics and cationic peptides can provide synergistic activity
CC against a microorganism that is tolerant, inherently resistant, or has
CC acquired resistance to an antibiotic agent. They can be used for killing
CC e.g. bacteria, fungi, parasites and viruses.
XX
S0 Sequence 13 AA:

Query Match 100.0%; Score 91; DB 19; Length 13;
Best Local Similarity 100.0%; Pred. NO: 7.2e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKKPMPWMPRRK 13
DB 1 ILKKPMPWMPRRK 13

RESULT 4
AAW71690
ID AAW71690 standard; Peptide: 13 AA.
XX
AC AAW71690;
XX
DT 11-JAN-1999 (first entry)
XX
DE Cationic peptide MB111 (MW 1879).
XX
KM MB111; cationic peptide; plasmid pK1; small cryptic plasmid;
KW replication; RepA; vector; RAMP.
XX
OS Synthetic.
XX
PN WO9841636-A2.
XX
PD 24-SEP-1998.
XX
PE 16-MAR-1998; 98MO-CA00214.
XX
PR 14-MAR-1997; 97US-0040722.
XX
PA (BURI/) BURIAN J.
PA (KAYW/) KAY W W.
XX
PI Burian J, Kay WW;
XX
XX WPI: 1998-531571/45.
XX
PT Increasing plasmid copy number in a cell with the repA gene product
PT - and an small cryptic plasmid ori sequence, useful for high level
PT expression of e.g. cytokines, antigens or therapeutic proteins
XX
XX Example 13; Page 54; 82pp; English.
XX
CC MB111 is a small (mol. wt. 1879) cationic peptide. DNA encoding
CC MB111 has been incorporated into vector pK1h-B1, in which the
CC replication leader (R21) sequence of RepA (see also AAW71686) is
CC joined to 2 Hpro peptides (see also AAW71692), to provide a
CC vector for expression of MB111 in host cells. The invention

CC provides controlled replication plasmid vectors (RAMP vectors)
CC comprising a replication origin of a small cryptic plasmid and a
CC gene encoding RepA. The vectors can reach very high levels of
CC plasmid replication, but are not lethal to the host cell, and can
CC be used to direct the high level expression of e.g. cytokines,
CC antigens and therapeutic proteins.
XX
S0 Sequence 13 AA:

Query Match 100.0%; Score 91; DB 19; Length 13;
Best Local Similarity 100.0%; Pred. NO: 7.2e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKKPMPWMPRRK 13
DB 1 ILKKPMPWMPRRK 13

RESULT 5
AA94495
ID AA94495 standard; Peptide: 13 AA.
XX
AC AA94495;
XX
DT 20-SEP-2000 (first entry)
XX
DE MB1-11 peptide derived from indolicidin.
XX
KM Cellulose binding domain; CBD; cationic peptide;
KM MB1-11; indolicidin; bovine.
XX
OS Bos taurus.
XX
PN WO200031279-A2.
XX
PD 02-JUN-2000.
XX
PE 19-NOV-1999; 99MO-CA01107.
XX
PR 20-NOV-1998; 98US-0109218.
XX
PA (MICR-) MICROLOGIX BIOTECH INC.
XX
PI Burian J, Bartfeld D;
XX
XX WPI: 2000-400086/34.
XX
DR Multi-domain fusion protein expression cassette used for high yield
XX stable production of foreign peptide gene products -
XX
XX Disclosure: Page 24; 73pp; English.
XX
XX A novel method allows the efficient production of cationic peptides in
XX recombinant host cells. The method involves construction of a
XX multi-domain fusion protein expression cassette comprising a promoter and
XX a nucleic acid molecule expressed as an insoluble protein. The inclusion
XX of anionic peptide sequences in the linker sequences neutralises the
XX positive charge of the cationic peptide so that the charge of the
XX fusion protein is controlled. This cassette allows high yield, stable
XX production of the cationic peptide. Cationic peptides such as
XX bovine indolicidin may be used as antimicrobial agents. The present
XX sequence is the MB1-11 peptide. MB1-11 is a cationic peptide derived
XX from modifications of indolicidin.
XX
S0 Sequence 13 AA:

Query Match 100.0%; Score 91; DB 21; Length 13;
Best Local Similarity 100.0%; Pred. NO: 7.2e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKKPMPWMPRRK 13

```
Db 1 ilkkwpwrk 13
```

RESULT	6
AA92795	
ID	AA92795 standard; peptide; 13 AA

AC AAY92795;

DT 29-AUG-2000 (first entry)

DE Indolicidin analogue, CP-11.

KW Magainin; antimicrobial; transgenic plant; protease degradation; Rev4
KW Indolicidin; protein production; reverse peptide.
....

Synthetic

PN WO200026344-A1.

PD 11-MAY-2000.

PF 29-OCT-1999; 99WO-US25561.

PR	30-OCT-1998;	98US-0106373
PR	03-NOV-1998;	98US-0106537

PA (INTE-) INTERLINK BIOTECHNOLOGIES LLC.
PA (KENT) UNIV KENTUCKY RES FOUND.

PI Everett NP, Li Q, Lawrence C, Davies MH;

DR WPI; 2000-365597/31.

PT Polypeptides for reducing proteolytic degradation of proteins
PT administered to, or produced by a plant comprise indolicin or its
PT functional equivalents

PS Disclosure; Page 4; 50pp; English

Indolicidin is a potent antimicrobial tridecapptide, originally purified from cytoplasmic granules of bovine neutrophils. Cp-11 is an analogue, which has better activity against *E. coli*, *Pseudomonas aeruginosa* and *Candida albicans*, but reduced activity against *Staphylococcus aureus*. A reverse peptide, Rev4 (AAW92796) of indolicidin was found to have increased stability against plant protease degradation. Expression of antimicrobial peptides in transgenic plants suffers a major limitation in that the foreign peptides are susceptible to rapid degradation by proteases. The invention concerns reducing the extent of protease degradation of a protein applied to, or produced by a plant by administering indolicidin, Rev4 or a functional equivalent to the plant. Transgenic plants expressing indolicidin and Rev4 are useful for production of the antimicrobial peptides. Compositions containing indolicidin and Rev4 are also useful for production of agronomically important proteins in plants.

Sequence 13 AA;

Query Match	100.0%	Score 91	.DB 21	Length 13
Best Local Similarity	100.0%	Pred. No.	7.2e-07	
Matches 13, Conservative	0	Mismatches	0	Indels 0
				Gaps 0

QY 1 ILKKWPWPWRK 13

Db 1 ilkkwpwrk 13

RESULT	7
AAV91773	
ID	AAV91773 standard; Peptide; 13 AA
XX	

AC	AA91773;
XX	
DT	06-JUN-2000 (first entry)

DE Amino acid sequence of cationic peptide MBI 11.

KW Cationic peptide; tumour; pharmaceutical composition; cancer; treatment.
KW leukaemia; polyalkylene-modified; APO; lymphoma; multiple myeloma;
KW breast; lung; ovary; cervix; uterus; prostate; liver; colon;
KW multidrug resistance.

OS Synthetic.

PN W09965506-A2.

PD 23-DEC-1999
XY

PF 14-JUN-1999; 99WO-CA005552
VY

PR 12-JUN-1998; 98US-0096541.

PA (MICR-) MICROLOGIX BIOTECH INC.
XX
XX

PI Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP,
XX
DR WPI, 2000-223549/19.

Novel pharmaceutical composition containing optionally activated
polyoxalylalkylene-modified cationic peptides, useful for treating tumours

PS Disclosure; Page 14; 94pp; English.

CC This sequence represents a cationic peptide amino acid sequence, which
CC can be used in the pharmaceutical composition of the invention. The
CC invention relates to a pharmaceutical composition containing at least one
CC activated polyoxalkylene (APO)-modified cationic peptide. The
CC modification of peptides with APO increases their activity against tumour
CC cells, including those with a multidrug resistant phenotype. The
CC pharmaceutical composition can be used to treat tumours, specifically
CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
CC cervix, uterus, skin, prostate, liver and colon.

SQ Sequence 13 AA;

Query Match	100.0%	Score 91;	DB 21;	Length 13;
Best Local Similarity	100.0%	Pred. No. 7	2e-07;	
Matches 13; Conservative	0;	Mismatches	0;	Indels 0; Gaps

QY	1	ILKKWPWPRK	13

Db 1 ilkkwrwrirk 13

RESULT	8
AAV91774	
ID	AAV91774 standard; Peptide; 13 AA

AC AAY91774

DT 06-JUN-2000 (first entry)

DE Amino acid sequence of cationic peptide MBI 11CN

KM Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
 KM leukaemia; polyoxalkylene-modified; APO; lymphoma; multiple myeloma;
 KM breast; lung; ovary; cervix; uterus; prostate; liver; colon;
 KM multidrug resistance.

OS	Synthetic
----	-----------

PN W09965506-A2

XX 23-DEC-1999.
 PD 14-JUN-1999; 99WO-CA00552.
 XX 12-JUN-1998; 98US-0096541.
 XX (MICR-) MICROLOGIX BIOTECH INC.
 PA Friedland HD, Krieger TJ, Taylor R, Erfile D, Fraser JR, West MHP;
 PI WPI; 2000-223549/19.
 XX Novel pharmaceutical composition containing optionally activated
 PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
 PS
 XX Example 3; Page 14; 94pp; English.
 CC This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxyalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.
 XX
 SQ Sequence 13 AA;
 Query Match *100.0%; Score 91; DB 21; Length 13;
 Best Local Similarity 100.0%; Pred. No. 7.2e-07;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ILKKPMPMPMRK 13
 Db 1 ILKKPMPMPMRK 13
 RESULT 9
 AAY91818
 ID AAY91818 standard; Peptide; 13 AA.
 XX AAY91818;
 AC
 XX 06-JUN-2000 (first entry).
 DT
 XX Amino acid sequence of cationic peptide MBI 11E1CN.
 DE
 XX Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
 KW leukaemia; polyoxyalkylene-modified; APO; lymphoma; multiple myeloma;
 KW breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
 KW multidrug resistance.
 XX
 XX Synthetic.
 OS
 XX WO9965506-A2.
 PN
 XX 23-DEC-1999.
 PD
 XX 14-JUN-1999; 99WO-CA00552.
 PF
 XX 12-JUN-1998; 98US-0096541.
 PR
 XX (MICR-) MICROLOGIX BIOTECH INC.
 PA Friedland HD, Krieger TJ, Taylor R, Erfile D, Fraser JR, West MHP;
 PI WPI; 2000-223549/19.
 XX Novel pharmaceutical composition containing optionally activated

PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
 PS
 XX Disclosure; Page 15; 94pp; English.
 CC This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxyalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.
 XX
 SQ Sequence 13 AA;
 Query Match 100.0%; Score 91; DB 21; Length 13;
 Best Local Similarity 100.0%; Pred. No. 7.2e-07;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ILKKPMPMPMRK 13
 Db 1 ILKKPMPMPMRK 13
 RESULT 10
 AAY91819
 ID AAY91819 standard; Peptide; 13 AA.
 XX AAY91819;
 AC
 XX 06-JUN-2000 (first entry)
 DT
 XX Amino acid sequence of cationic peptide MBI 11E2CN.
 DE
 XX Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
 KW leukaemia; polyoxyalkylene-modified; APO; lymphoma; multiple myeloma;
 KW breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
 KW multidrug resistance.
 XX
 XX Synthetic.
 OS
 XX WO9965506-A2.
 PN
 XX 23-DEC-1999.
 PD
 XX 14-JUN-1999; 99WO-CA00552.
 PF
 XX 12-JUN-1998; 98US-0096541.
 PR
 XX (MICR-) MICROLOGIX BIOTECH INC.
 PA Friedland HD, Krieger TJ, Taylor R, Erfile D, Fraser JR, West MHP;
 PI WPI; 2000-223549/19.
 XX Novel pharmaceutical composition containing optionally activated
 PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
 PS
 XX Disclosure; Page 15; 94pp; English.
 CC This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxyalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.

XX Sequence 13 AA;

Query Match
Best Local Similarity 100.0%; Score 91; DB 21; Length 13;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWPRRK 13
Db 1 ILKKPMPWPRRK 13

RESULT 11

AA91820
ID AAY91820 standard; Peptide; 13 AA.

AC AAY91820;

DT 06-JUN-2000 (first entry);

DE Amino acid sequence of cationic peptide MBI 11E3CN.

KW Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
KW leukaemia; polyoxyalkylene-modified; APO; lymphoma; multiple myeloma;
KW breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;

XX multidrug resistance.

OS Synthetic.

PN WO965506-A2.

PD 23-DEC-1999.

PF 14-JUN-1999; 99WO-CA00552.

PR 12-JUN-1998; 98US-0096541.

PI (MICR-) MICROLOGIX BIOTECH INC.

PI Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP;

DR WPI; 2000-223549/19.

PT Novel pharmaceutical composition containing optionally activated
PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
PS Claim 1; Page 15; 94pp; English.

CC This sequence represents a cationic peptide amino acid sequence, which
CC can be used in the pharmaceutical composition of the invention. The
CC invention relates to a pharmaceutical composition containing at least one
CC activated polyoxyalkylene (APO)-modified cationic peptide. The
CC modification of peptides with APO increases their activity against tumour
CC cells, including those with a multidrug resistant phenotype. The
CC pharmaceutical composition can be used to treat tumours, specifically
CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
CC cervix, uterus, skin, prostate, liver and colon.

CC Sequence 13 AA;

Query Match
Best Local Similarity 100.0%; Score 91; DB 21; Length 13;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWPRRK 13
Db 1 ILKKPMPWPRRK 13

RESULT 12

AA924583
ID AAY24583 standard; peptide; 14 AA.

AC AAY24583;

DT 18-AUG-1999 (first entry)

DE Indolicidin analogue #35.

KW Indolicidin; bacterial infection; photo-oxidised solubiliser;
KW antimicrobial; antibiotic; antitumour; antitumour; surface disinfectant;
KW additive; shampoo; soap; insecticide; herbicide; preservative;
KW food; technical material.

OS Synthetic.

PN WO9807745-A2.

PD 26-FEB-1998.

PF 21-AUG-1997; 97WO-US14779.

PR 13-JAN-1997; 97US-0034949.

PR 21-AUG-1996; 96US-0024754.

PI (MICR-) MICROLOGIX BIOTECH INC.

PI Erfle D, Fraser JR, Krieger TJ, Taylor R, West MHP;

DR WPI; 1998-169090/15.

PT New indolicidin analogues with antimicrobial activity and related
PT nucleic acid vectors, transformed cells and antibodies, also
PT conjugates with polyoxyalkylene glycol and fatty acid to reduce
PT toxicity, useful therapeutically, as disinfectants etc.

PS Claim 13; Page 89; 129pp; English.

CC AAY24549 to AAY24615 represent indolicidin analogues of formulae
CC (I)-(VIII) containing up to 25 amino acids (aa): R₁XX₁XB₁(I), B₁XX₁XB₁
CC (II), B₁B₁XX₁XB₁(III), B₁XX₁XB₁B₁(AA)n₁ILB₁B₁AGS (IV), B₁XX₁XB₁B₁(AA)n₁m
CC (V), LBB₁XX₁XB₁XX₁XB₁ (VI), LK₁XX₁XB₁XX₁XB₁ (VII) and B₁XX₁XB₁XX₁XB₁ (VIII).
CC Where Z = P or V; X = hydrophobic residue, preferably W; B = basic aa;
CC preferably R or K; AA = any aa; n = 0 or 1; in (II), at least 1 Z = V;
CC in (VIII) at least 2 X = F or Y. The analogues are used to treat
CC infections caused by bacteria (Gram positive or negative, or anaerobic);
CC fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
CC trematodes) or viruses. Typical of very many pathogens that can be
CC controlled are Leishmania, Trypanosoma, Ascaris lumbricoides, Fasciola
CC hepatica, Klebsiella pneumoniae, Bordetella pertussis, Staphylococcus
CC aureus, Listeria, Clostridium, rotavirus and papilloma virus. Compounds
CC also derived from the analogues may be used similarly; the compounds may
CC may be used therapeutically or to coat medical devices; also they are
CC useful as surface disinfectants, as additives to shampoo or soaps, as
CC insecticides or herbicides, or as preservatives for foods or soaps, as
CC materials. The analogues are administered by injection, lavage, orally
CC or topically, generally at 0.1-50 mg/kg. These analogues have a broader
CC spectrum of activity than indolicidin and modification as compounds
CC reduces their toxicity.

CC Sequence 14 AA;

Query Match
Best Local Similarity 100.0%; Score 91; DB 19; Length 14;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWPRRK 13
Db 1 ILKKPMPWPRRK 13

RESULT 13
AA91811
ID AAY91811 standard; Peptide: 14 AA.
XX
AC AAY91811;
XX
DT 06-JUN-2000 (first entry)
XX
DE Amino acid sequence of cationic peptide MBI 11D11H.
XX
KW Cationic peptide: tumour; pharmaceutical composition: cancer; treatment;
KW leukaemia; polyoxalkylene-modified; APO; lymphoma; multiple myeloma;
KW breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
KW multidrug resistance.
XX
OS Synthetic.
XX
PN WO9965506-A2.
XX
PD 23-DEC-1999.
XX
PE 14-JUN-1999; 99WO-CA00552.
XX
PR 12-JUN-1998; 98US-0096541.
XX
PA (MICR-) MICROLOGIX BIOTECH INC.
XX
PI Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP;
XX
DR WPI: 2000-223549/19.
XX
PT Novel pharmaceutical composition containing optionally activated
PT polyoxalkylene-modified cationic peptides, useful for treating tumours
PT
XX
PS Disclosure; Page 15; 94pp; English.
XX
SS This sequence represents a cationic peptide amino acid sequence, which
CC can be used in the pharmaceutical composition of the invention. The
CC invention relates to a pharmaceutical composition containing at least one
CC activated polyoxalkylene (APO)-modified cationic peptide. The
CC modification of peptides with APO increases their activity against tumour
CC cells, including those with a multidrug resistant phenotype. The
CC pharmaceutical composition can be used to treat tumours, specifically
CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
CC cervix, uterus, skin, prostate, liver and colon.
CC
XX
SQ Sequence 14 AA:

Query Match 100.0%; Score 91; DB 21; Length 14;
Best Local Similarity 100.0%; Pred. No. 7.8e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKKPMPMPRRK 13
DB 1 ILKKPMPMPRRK 13

RESULT 14
AAY24582
ID AAY24582 standard; peptide: 21 AA.
XX
AC AAY24582;
XX
DT 18-AUG-1999 (first entry)
XX
DE Indolicidin analogue #34.
XX
KW Indolicidin; bacterial infection; photo-oxidised solubiliser;
KW antimicrobial; antibiotic; antiarrhythmic; surface disinfectant;
KW additive; shampoo; soap; insecticide; herbicide; preservative;
KW food; technical material.

XX
OS Synthetic.
XX
PN WO9807745-A2.
XX
PD 26-FEB-1998.
XX
PE 21-AUG-1997; 97WO-US14779.
XX
PR 13-JAN-1997; 97US-0034949.
PR 21-AUG-1996; 96US-0024754.
XX
PA (MICR-) MICROLOGIX BIOTECH INC.
XX
PI Erfle D, Fraser JR, Krieger TJ, Taylor R, West MH;
XX
DR WPI: 1998-169090/15.
XX
PT New indolicidin analogues with antimicrobial activity and related
PT nucleic acid - vectors, transformed cells and antibodies, also
PT conjugates with polyoxalkylene glycol and fatty acid to reduce
PT toxicity, useful therapeutically, as disinfectants etc.
XX
PS Claim 13; Page 89; 129pp; English.
XX
SS AAY24549 to AAY24615 represent indolicidin analogues of formulae
CC (I)-(VIII) containing up to 25 amino acids (aa): RXXXXZXB (I), BXXXXZXB
CC (II), BBXXZXXZXB (III), BXXZXXZXBn(AA)nMLBBnGS (IV), BXXZXXZXB(AA)nM
CC (V), LBBnXZnXXZnXRK (VI), LKnXXZXXZRRK (VII) and BXXZXXZXBn (VIII).
CC Where Z = P or V; X = hydrophobic residue, preferably W; B = basic aa;
CC preferably R or K; AA = any aa; n = 0 or 1; In (II), at least 1 Z = V;
CC in (VIII) at least 2 X = F or Y. The analogues are used to treat
CC infections caused by bacteria (Gram positive or negative, or anaerobic);
CC fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
CC trematodes) or viruses. Typical of very many pathogens that can be
CC controlled are Leishmania, Trypanosoma, Ascaris lumbricoides, Fasciola
CC hepatica, Klebsiella pneumoniae, Bordetella pertussis, Staphylococcus
CC aureus, Listeria, Clostridium, rotavirus and papilloma virus. Compounds
CC derived from the analogues may be used similarly: the compounds may
CC also be prepared from antibiotics or antiarrhythmic agents. The analogues
CC may be used therapeutically or to coat medical devices; also they are
CC useful as surface disinfectants, as additives to shampoo or soaps, as
CC insecticides or herbicides, or as preservatives for foods and technical
CC materials. The analogues are administered by injection, lavage, orally
CC or topically, generally at 0.1-50 mg/kg. These analogues have a broader
CC spectrum of activity than indolicidin and modification as compounds
CC reduces their toxicity.
XX
SQ Sequence 21 AA:

Query Match 100.0%; Score 91; DB 19; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.2e-06;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKKPMPMPRRK 13
DB 1 ILKKPMPMPRRK 13

RESULT 15
AAY91806
ID AAY91806 standard; Peptide: 21 AA.
XX
AC AAY91806;
XX
DT 06-JUN-2000 (first entry)
XX
DE Amino acid sequence of cationic peptide MBI 11D4CN.
XX
KW Cationic peptide: tumour; pharmaceutical composition: cancer; treatment;
KW leukaemia; polyoxalkylene-modified; APO; lymphoma; multiple myeloma;
KW breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;

KM multidrug resistance.
 XX
 OS Synthetic.
 XX
 PN WO9965506-A2.
 XX
 PD 23-DEC-1999.
 XX
 PF 14-JUN-1999; 99WO-CA00552.
 XX
 PR 12-JUN-1998; 98US-0096541.
 XX
 PA (MICR-) MICROLOGIX BIOTECH INC.
 XX
 PI Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP;
 XX
 DR WPI; 2000-223549/19.
 XX
 PT Novel pharmaceutical composition containing optionally activated
 PT polyoxalkylene-modified cationic peptides, useful for treating tumours
 PT
 PS Disclosure; Page 15; 94pp; English.
 XX
 CC This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.
 CC
 SQ Sequence 21 AA;

Query Match 100.0%; Score 91; DB 21; Length 21;
 Best Local Similarity 100.0%; Pred. No. 1.2e-06;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 ILKKPMPMPRRK 13
 |||||
 Db 1 ilkkpmpmprrk 13

Search completed: January 4, 2002, 08:40:26
 Job time: 109 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:40:57 ; Search time 24.75 Seconds
(without alignments)
11.820 Million cell updates/sec

Title: US-09-444-281-35

Perfect score: 91

Sequence: 1 ILKKWPMWRRK 13

Scoring table: BLOSUM62

Searched: Gapop 10.0 , Gapext 0.5

Total number of hits satisfying chosen parameters: 212252

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Database :

Issued_Patents_AA:*
1: /cgn2_6/ptodata/2/1aa/5A.COMB.pep:*
2: /cgn2_6/ptodata/2/1aa/5B.COMB.pep:*
3: /cgn2_6/ptodata/2/1aa/6A.COMB.pep:*
4: /cgn2_6/ptodata/2/1aa/6B.COMB.pep:*
5: /cgn2_6/ptodata/2/1aa/PCTUS.COMB.pep:*
6: /cgn2_6/ptodata/2/1aa/backfiles1.pep:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	91	100.0	13	4	US-08-915-314-30 Sequence 30, App1
2	91	100.0	13	4	US-08-915-314-62 Sequence 62, App1
3	91	100.0	13	4	US-08-915-314-63 Sequence 63, App1
4	91	100.0	13	4	US-08-915-314-64 Sequence 64, App1
5	91	100.0	13	4	US-09-042-071-36 Sequence 36, App1
6	91	100.0	14	4	US-08-915-314-57 Sequence 57, App1
7	91	100.0	21	4	US-08-915-314-54 Sequence 54, App1
8	87	95.6	12	4	US-08-915-314-52 Sequence 52, App1
9	86	94.5	12	4	US-08-915-314-74 Sequence 74, App1
10	86	94.5	12	4	US-08-702-054B-5 Sequence 5, App1
11	86	94.5	13	4	US-08-915-314-58 Sequence 58, App1
12	86	94.5	14	4	US-08-915-314-59 Sequence 59, App1
13	86	94.5	15	4	US-08-702-054B-40 Sequence 40, App1
14	86	94.5	20	4	US-08-915-314-55 Sequence 55, App1
15	86	94.5	21	4	US-08-915-314-56 Sequence 56, App1
16	85	93.4	12	4	US-08-915-314-69 Sequence 69, App1
17	85	93.4	13	4	US-08-915-314-38 Sequence 38, App1
18	85	93.4	13	4	US-08-915-314-45 Sequence 45, App1
19	85	93.4	13	4	US-08-702-054B-1 Sequence 1, App1
20	85	93.4	13	4	US-08-702-054B-17 Sequence 17, App1
21	85	93.4	13	4	US-08-702-054B-32 Sequence 32, App1
22	83	91.2	12	4	US-08-915-314-24 Sequence 24, App1
23	83	91.2	13	4	US-08-915-314-49 Sequence 49, App1
24	83	91.2	13	4	US-08-915-314-50 Sequence 50, App1
25	83	91.2	13	4	US-08-915-314-51 Sequence 51, App1
26	83	91.2	13	4	US-08-702-054B-30 Sequence 30, App1
27	83	91.2	13	4	US-08-702-054B-31 Sequence 31, App1

28	83	91.2	13	4	US-08-702-054B-34 Sequence 34, App1
29	83	91.2	13	4	US-08-702-054B-35 Sequence 35, App1
30	82	90.1	13	4	US-08-915-314-25 Sequence 25, App1
31	82	90.1	13	4	US-08-915-314-66 Sequence 66, App1
32	82	90.1	13	4	US-08-915-314-67 Sequence 67, App1
33	82	90.1	13	4	US-08-702-054B-33 Sequence 33, App1
34	81	89.0	11	4	US-08-915-314-75 Sequence 75, App1
35	81	89.0	15	4	US-08-702-054B-39 Sequence 39, App1
36	80	87.9	14	4	US-08-702-054B-18 Sequence 18, App1
37	80	87.9	15	4	US-08-702-054B-41 Sequence 41, App1
38	80	87.9	16	4	US-08-702-054B-2 Sequence 2, App1
39	79.5	87.4	16	4	US-08-702-054B-38 Sequence 38, App1
40	79	86.8	17	4	US-08-702-054B-42 Sequence 42, App1
41	78	85.7	11	4	US-08-915-314-28 Sequence 28, App1
42	78	85.7	12	4	US-08-915-314-40 Sequence 40, App1
43	77	84.6	12	4	US-08-915-314-39 Sequence 39, App1
44	77	84.6	12	4	US-08-702-054B-27 Sequence 27, App1
45	76	83.5	12	4	US-08-915-314-77 Sequence 77, App1

ALIGNMENTS

RESULT 1
US-08-915-314-30
Sequence 30, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erle, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED AND BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
City: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 30:
SEQUENCE CHARACTERISTICS:
LENGTH: 13 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-30

Query Match 100.0%; Score 91; DB 4; Length 13;
Best Local Similarity 100.0%; Pred. No. 5.4e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ILKKWPMWRRK 13

Db 1 ILKKPMPWPRRK 13
RESULT 2
US-08-915-314-62.
Sequence 62, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915, 314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 62:
SEQUENCE CHARACTERISTICS:
LENGTH: 13 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
FEATURE:
NAME/KEY: Modified-site
LOCATION: 1
OTHER INFORMATION: /note="D-Form of Isoleucine"
US-08-915-314-62
Query Match 100.0%; Score 91; DB 4; Length 13;
Best Local Similarity 100.0%; Pred. No. 5.4e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 1 ILKKPMPWPRRK 13
Db 1 ILKKPMPWPRRK 13
RESULT 3
US-08-915-314-63
Sequence 63, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:

TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915, 314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 63:
SEQUENCE CHARACTERISTICS:
LENGTH: 13 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
FEATURE:
NAME/KEY: Modified-site
LOCATION: 13
OTHER INFORMATION: /note="D-Form of Lysine"
US-08-915-314-63
Query Match 100.0%; Score 91; DB 4; Length 13;
Best Local Similarity 100.0%; Pred. No. 5.4e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 1 ILKKPMPWPRRK 13
Db 1 ILKKPMPWPRRK 13
RESULT 4
US-08-915-314-64
Sequence 64, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/915:314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 64:
SEQUENCE CHARACTERISTICS:
LENGTH: 13 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
FEATURE:
NAME/KEY: Modified-site
LOCATION: 1
OTHER INFORMATION: /note="D-Form of Isoleucine"
FEATURE:
NAME/KEY: Modified-site
LOCATION: 13
OTHER INFORMATION: /note="D-Form of Lysine"
US-08-915-314-64

Query Match 100.0%; Score 91; DB 4; Length 13;
Best Local Similarity 100.0%; Pred. No. 5.4e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWMPRRK 13
Db 1 ILKKPMPWMPRRK 13

RESULT 5
US-09-042-071-36
Sequence 36, Application US/09042071
Patent No. 6294372
GENERAL INFORMATION:
APPLICANT: Burian, Jan
APPLICANT: Kay, William W.
TITLE OF INVENTION: REPLICATION GENES AND GENE PRODUCTS FROM
TITLE OF INVENTION: SMALL CRYPTIC PLASMIDS AND METHODS FOR CONSTRUCTING
TITLE OF INVENTION: CONTROLLED-REPLICATION PLASMID VECTORS
NUMBER OF SEQUENCES: 52
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/042.071
FILING DATE: 13-MAR-1998
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Mcmasters, David D.
REGISTRATION NUMBER: 33,963
REFERENCE/DOCKET NUMBER: 660081.407
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 36:
SEQUENCE CHARACTERISTICS:
LENGTH: 13 amino acids

TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
US-09-042-071-36

Query Match 100.0%; Score 91; DB 4; Length 13;
Best Local Similarity 100.0%; Pred. No. 5.4e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWMPRRK 13
Db 1 ILKKPMPWMPRRK 13

RESULT 6
US-08-915-314-57
Sequence 57, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfle, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915.314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 57:
SEQUENCE CHARACTERISTICS:
LENGTH: 14 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-57

Query Match 100.0%; Score 91; DB 4; Length 14;
Best Local Similarity 100.0%; Pred. No. 5.8e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWMPRRK 13
Db 1 ILKKPMPWMPRRK 13

RESULT 7
US-08-915-314-54
Sequence 54, Application US/08915314
Patent No. 6180604

GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESSES:
ADDRESSSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 54:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-54

Query Match 100.0%; Score 91; DB 4; Length 21;
Best Local Similarity 100.0%; Pred. No. 8.5e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKKPMPWPRRK 13
Db 1 ILKKPMPWPRRK 13

RESULT 8
US-08-915-314-52
Sequence 52, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESSES:
ADDRESSSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 52:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-52

Query Match 95.6%; Score 87; DB 4; Length 12;
Best Local Similarity 100.0%; Pred. No. 1.8e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 LKKPMPWPRRK 13
Db 1 LKKPMPWPRRK 12

RESULT 9
US-08-915-314-74
Sequence 74, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESSES:
ADDRESSSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 74:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear

US-08-915-314-74

Query Match 94.5%; Score 86; DB 4; Length 12;

Best Local Similarity 100.0%; Pred. No. 2.4e-06;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWPMRR 12
| | | | | | | | | | | | | |
Db 1 ILKKPMPWPMRR 12

RESULT 10

US-08-702-054B-5
Sequence 5, Application US/08702054B
Patent No. 6191254

GENERAL INFORMATION:

APPLICANT: Falls, Timothy J.

APPLICANT: Hancock, Robert E. W.

APPLICANT: Gough, Monisha

TITLE OF INVENTION: ANTIMICROBIAL CATIONIC PEPTIDES

TITLE OF INVENTION: AND METHODS OF SCREENING FOR THE SAME

NUMBER OF SEQUENCES: 44

CORRESPONDENCE ADDRESS:

ADDRESSEE: Fish & Richardson P.C.

STREET: 4225 Executive Square, Suite 1400

CITY: La Jolla

STATE: CA

COUNTRY: USA

ZIP: 92037

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

COMPUTER: IBM Compatible

OPERATING SYSTEM: Windows 95

SOFTWARE: FastSeq for Windows Version 2.0b

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/702.054B

FILING DATE: 23-AUG-1996

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 60/002.687

FILING DATE: 23-AUG-1995

ATTORNEY/AGENT INFORMATION:

NAME: Halle, Lisa A.

REGISTRATION NUMBER: 38,347

REFERENCE/DOCKET NUMBER: 07420/013001

TELEPHONE: 619/678-5070

TELEFAX: 619/678-5099

INFORMATION FOR SEQ ID NO: 5:

SEQUENCE CHARACTERISTICS:

LENGTH: 12 amino acids

TYPE: amino acid

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-08-702-054B-5

Query Match 94.5%; Score 86; DB 4; Length 12;

Best Local Similarity 100.0%; Pred. No. 2.4e-06;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWPMRR 12
| | | | | | | | | | | | | |
Db 1 ILKKPMPWPMRR 12

RESULT 11

US-08-915-314-58

Sequence 58, Application US/08915314

Patent No. 6180604

GENERAL INFORMATION:

APPLICANT: Fraser, Janet R.

APPLICANT: West, Michael H.P.

APPLICANT: Krieger, Timothy J.

APPLICANT: Taylor, Robert

APPLICANT: Erfle, Douglas

TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING

TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN

NUMBER OF SEQUENCES: 90

CORRESPONDENCE ADDRESS:

ADDRESSEE: SEED and BERRY LLP

STREET: 6300 Columbia Center, 701 Fifth Avenue

CITY: Seattle

STATE: Washington

COUNTRY: USA

ZIP: 98104

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/915.314

FILING DATE: 20-AUG-1997

CLASSIFICATION: 424

ATTORNEY/AGENT INFORMATION:

NAME: No. 6180604tenburg Ph.D., Carol

REGISTRATION NUMBER: 39,317

REFERENCE/DOCKET NUMBER: 660081.405

TELEPHONE: (206) 622-4900

TELEFAX: (206) 682-6031

INFORMATION FOR SEQ ID NO: 58:

SEQUENCE CHARACTERISTICS:

LENGTH: 13 amino acids

TYPE: amino acid

STRANDEDNESS:

TOPOLOGY: linear

US-08-915-314-58

Query Match 94.5%; Score 86; DB 4; Length 13;

Best Local Similarity 100.0%; Pred. No. 2.0e-06;

Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWPMRR 12
| | | | | | | | | | | | | |
Db 1 ILKKPMPWPMRR 12

RESULT 12

US-08-915-314-59

Sequence 59, Application US/08915314

Patent No. 6180604

GENERAL INFORMATION:

APPLICANT: Fraser, Janet R.

APPLICANT: West, Michael H.P.

APPLICANT: Krieger, Timothy J.

APPLICANT: Taylor, Robert

APPLICANT: Erfle, Douglas

TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING

TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN

NUMBER OF SEQUENCES: 90

CORRESPONDENCE ADDRESS:

ADDRESSEE: SEED and BERRY LLP

STREET: 6300 Columbia Center, 701 Fifth Avenue

CITY: Seattle

STATE: Washington

COUNTRY: USA

ZIP: 98104

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604 Leuburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 59:
SEQUENCE CHARACTERISTICS:
LENGTH: 14 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-59

Query Match 94.5%; Score 86; DB 4; Length 14;
Best Local Similarity 100.0%; Pred. No. 2.8e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPWPMPWR 12
DB 1 ILKKPWPMPWR 12

RESULT 13
US-08-702-054B-40
Sequence 40, Application US/08702054B
Patent No. 6191254
GENERAL INFORMATION:
APPLICANT: Falls, Timothy J.
APPLICANT: Hancock, Robert E. W.
TITLE OF INVENTION: ANTIMICROBIAL CATIONIC PEPTIDES
TITLE OF INVENTION: AND METHODS OF SCREENING FOR THE SAME
NUMBER OF SEQUENCES: 44
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson P.C.
STREET: 4225 Executive Square, Suite 1400
CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: FASTSEQ for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/702,054B
FILING DATE: 23-AUG-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/002,687
FILING DATE: 23-AUG-1995
ATTORNEY/AGENT INFORMATION:
NAME: Haile, Lisa A.
REGISTRATION NUMBER: 38,347
REFERENCE/DOCKET NUMBER: 07420/013001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 619/678-5070
TELEFAX: 619/678-5099
INFORMATION FOR SEQ ID NO: 40:
SEQUENCE CHARACTERISTICS:
LENGTH: 15 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-702-054B-40

Query Match 94.5%; Score 86; DB 4; Length 15;
Best Local Similarity 100.0%; Pred. No. 3e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPWPMPWR 12
DB 1 ILKKPWPMPWR 12

RESULT 14
US-08-915-314-55
Sequence 55, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104

COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604 Leuburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 55:
SEQUENCE CHARACTERISTICS:
LENGTH: 20 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-55

Query Match 94.5%; Score 86; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.9e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPWPMPWR 12
DB 1 ILKKPWPMPWR 12

RESULT 15
US-08-915-314-56
Sequence 56, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfile, Douglas

TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
 TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
 NUMBER OF SEQUENCES: 90
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: SEED AND BERRY LLP
 STREET: 6300 Columbia Center, 701 Fifth Avenue
 CITY: Seattle
 STATE: Washington
 COUNTRY: USA
 ZIP: 98104
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/915,314
 FILING DATE: 20-AUG-1997
 CLASSIFICATION: 424
 ATTORNEY/AGENT INFORMATION:
 NAME: No. 6180604tenburg Ph.D., Carol
 REGISTRATION NUMBER: 39,317
 REFERENCE/DOCKET NUMBER: 660081.405
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (206) 622-4900
 TELEFAX: (206) 682-6031
 INFORMATION FOR SEQ ID NO: 56:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 21 amino acids
 TYPE: amino acid
 STRANDEDNESS:
 TOPOLOGY: linear
 SS-08-915-314-56

Query Match Similarity	94.5%	score	86;	DB	4;	length	21;
Best Local Similarity	100.0%	Pred. No.	4	1e-06;			
Matches	12;	Conservative	0;	Mismatches	0;	Indels	0;
						Gaps	0;

Qy	1	ILKKWPWPWRR	12
Db	1	ILKKWPWPWRR	12

```

Search completed: January 4, 2002, 08:40:57
Job time: 140 sec

```

THIS PAGE BLANK (USPTO)

OM of: US-09-444-281-35 to: GenEmbl:* out_format: pfs
Date: Jan 4, 2002 10:55 AM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-2000 Compugen Ltd.

Command line parameters:

-MODEL-frame.p2n.model -DEV=xlp
-O/cg22.1/USPTO.spool/US09444281/runat_04012002_084143_16195/app-query.fasta_1.210
-DB-GenEmbl -QFMT=fastap -SUFFIX=range -GAPOP=12.000 -GAPEXT=4.000
-MINMATCH=0.100 -LOOPL=0.000 -LOOPEXT=0.000 -OGAPOP=4.500
-OGAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500 -XGAPEXT=6.000
-FGAPEXT=7.000 -YGAPEXT=10.000 -YGAPEXT=0.500 -DEL0P=6.000
-DELXT=7.000 -STAR=1 -MATRIX=blotsum62 -TRANS=human0.cdi
-LIST=45 -DOCALLIGN=200 -THR_SCORE=pcr -THR_MAX=100 -THR_MIN=0
-ALIGN=15 -MODE=LOCAL -OUTFMT=pfs -NORM=ext -MINLEN=0
-MAXLEN=2000000000 -USER=US09444281_GCGN1_15145 -NCPU=6
-ICPU=3 -LONGLOG -NO_XLPHY -WAIT -THREDS=1

Search information block:

Query: US-09-444-281-35
Query length: 13
Database: GenEmbl:*
Database sequences: 1472140
Database length: 341344837
Search time (sec): 1728.450000

score list:

Sequence	Strd Orig	ZScore	EScore Len	Documentation
gb_om:BTINDLCD	73.00	158.82	1.43	550 X67340 B.taurus mRNA for indol
gb_pat:AX098418	73.00	11.25	13.65	6446 AX098418 Sequence 5 from Paten
gb_htg:AP003754	68.00	109.17	836.02	129052 AP003754 Oryza sativa chrom
gb_ro:MMU08210	66.00	132.63	41.24	2651 U08210 Mus musculus tncpola
gb_ro:AF289665	66.00	106.21	1.2e+03	107257 AF289665 Mus musculus E1F4H
gb_htg:AC091250	66.00	101.74	2.2e+03	200849 AC091250 Mus musculus chrom
gb_htg:AC090365	65.00	113.72	466.29	22780 AC090365 Drosophila melanog
gb_in:AC008316	65.00	101.18	2.3e+03	160817 AC008316 Drosophila melanog
gb_in:AC008315	65.00	100.50	2.5e+03	177028 AC008315 Drosophila melanog
gb_in:AE003684	65.00	98.96	3.1e+03	219579 AE003684 Drosophila melanog
gb_ro:WMA21XCA	64.00	114.42	443.17	19479 W22923 M.musculus alpha2 (1x)
gb_pr:AL162423	64.00	103.32	1.8e+03	88323 AL162423 Human DNA sequence
gb_htg:AL356097_0	64.00	101.76	2.2e+03	110000 AL356097 Homo sapiens chrom
gb_htg:AC072113	64.00	99.11	3.0e+03	159391 AC072113 Homo sapiens chrom
gb_htg:AL358473	64.00	97.94	3.5e+03	187445 AL358473 Homo sapiens chrom
gb_ro:AF210429	63.00	138.78	18.73	456 AF210429 Mus musculus group X
gb_ro:AF166097	63.00	132.90	39.85	1040 AF166097 Mus musculus group X
gb_ro:AF191309	63.00	124.85	111.95	3213 AF191309 Mus musculus zinc fir
gb_ro:RNY17153	63.00	120.98	183.83	5523 RNY17153 Rattus norvegicus mRNA
gb_ro:RATNCNHS	63.00	119.33	227.10	6957 M26643 Rat skeletal muscle vol
gb_in:CFR4A8	63.00	112.95	515.06	17015 Z61539 Caenorhabditis elegans
gb_da:AE004649	63.00	98.53	690.23	23425 AE004649 Pseudomonas aerugin
gb_ro:AC003063	63.00	95.77	3.3e+03	128143 AC003063 Mus musculus chrom
gb_ro:AC005817	63.00	95.77	4.7e+03	188707 AC005817 Mus musculus chrom
gb_htg:AC078895	63.00	94.95	5.2e+03	213204 AC078895 Mus musculus chrom
gb_htg:AC079043	63.00	94.85	5.2e+03	214723 AC079043 Mus musculus chrom
gb_pat:AX078941	62.00	130.84	51.92	1029 AX078941 Sequence 17 from Pat
gb_pl:ATPHB	61.50	116.30	199.31	3850 X17342 Arabidopsis thaliana ph
gb_pl:CELM01G5	61.50	120.65	322.37	37675 L09262 Arabidopsis thaliana p
gb_pl:AC005724	61.50	98.12	1.6e+03	36671 AC005724 Caenorhabditis eleg
gb_pl:AC003141	61.50	94.63	3.5e+03	141275 AC003141 Oryza sativa genom
gb_htg:AC013145	61.00	104.66	1.5e+03	29692 AC013145 Drosophila melanog
gb_in:DMR37P7	61.00	95.84	4.6e+03	102619 AL050231 Drosophila melanog
gb_htg:AC010475	61.00	94.44	5.5e+03	124927 AC010475 Homo sapiens chrom
gb_htg:AC092745	61.00	94.44	5.5e+03	125047 AC092745 Homo sapiens chrom
gb_pr:AC010485	61.00	94.41	5.6e+03	125439 AC010485 Homo sapiens chrom
gb_htg:AC080046	61.00	93.54	6.2e+03	141674 AC080046 Homo sapiens chrom
gb_htg:AP003910	61.00	93.44	6.3e+03	143549 AP003910 Oryza sativa chrom
gb_htg:AC083933	61.00	93.32	6.3e+03	143717 AC083933 Homo sapiens chrom
gb_pl:AP001366	61.00	93.32	6.4e+03	146081 AP001366 Oryza sativa genom

gb_htg:AC068010 - 61.00 92.42 7.2e+03 165652 | AC068010 Homo sapiens chr
gb_htg:AC063927 - 61.00 92.26 7.3e+03 169491 | AC063927 Homo sapiens chr
gb_htg:AC067797 - 61.00 91.99 7.6e+03 175940 | AC067797 Homo sapiens chr
gb_in:AE003417 - 61.00 87.84 1.3e+04 314661 | AE003417 Drosophila melan

seq_name: gb_om:BTINDLCD

seq_documentation_block:

LOCUS BTINDLCD 550 bp mRNA NAM 07-OCT-1992
DEFINITION B.taurus mRNA for indolcidin.
ACCESSION X67340
VERSION X67340.1 GI:462
KEYWORDS indolcidin.
SOURCE cow.
ORGANISM Bos taurus

REFERENCE 1 (bases 1 to 550)
1 (bases 1 to 550)
Del Sal G., Storici P., Schneider C., Romeo D. and Zanetti M.
Bovidae; Bovineae; Bos.
Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
Bovidae; Bovineae; Bos.

REFERENCE 2 (bases 1 to 550)
2 (bases 1 to 550)
Del Sal G.
Bovidae; Bovineae; Bos.
Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
Bovidae; Bovineae; Bos.

REFERENCE 3 (bases 1 to 550)
3 (bases 1 to 550)
Del Sal G.
Bovidae; Bovineae; Bos.
Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
Bovidae; Bovineae; Bos.

REFERENCE 4 (bases 1 to 550)
4 (bases 1 to 550)
Del Sal G.
Bovidae; Bovineae; Bos.
Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
Bovidae; Bovineae; Bos.

FEATURES

source

1..550
location/Qualifiers

/organism="Bos taurus"
/db_xref="taxon:9913"
/cell_line="Bone marrow cells"
13..99
13..447
/codon_start=1
/product="indolcidin prepropeptide"
/protein_id="CAA47755.1"
/db_xref="GI:463"
/translation="MOTQASLSIGRSLMLLLGLVPSASQAQSYRAVLRAVDO
LNESEANVRLLEDPPEKMDLSTRPVSFTVETVCPTTQOPARQCFKEKG
RIVOCVETVLDPSNDQFDLNCDELGVILPMKMPMPMRG"

sig_peptide

403..444
/product="indolcidin"
115 t

BASE COUNT

126 a 140 c 169 g 115 t

ORIGIN

126 a 140 c 169 g 115 t

alignment_scores:

Quality: 73.00 Length: 9
Ratio: 8.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-35 x BTINDLCD
Align seg 1/1 to: BTINDLCD from: 1 to: 550

4

12
|||||

415

441

seq_name: gb_pat:AX098418

415

seq_documentation_block:

6446 bp mRNA PAT 03-APR-2001

LOCUS AX098418

6446 bp mRNA PAT 03-APR-2001

DEFINITION

Sequence 5 from Patent WO0119969.

ACCESSION

AX098418

VERSION

AX098418.1 GI:13537710

KEYWORDS

Nicotiana benthamiana.

ORGANISM Nicotiana benthamiana, Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Asteridae; euasterids I; Solanales; Solanaceae; Nicotiana.

REFERENCE 1 (bases 1 to 6446)
Garger,S.J., Holtz,B.R., McCulloch,M.J. and Turner,T.H.
A process for isolating and purifying viruses, soluble proteins and peptides from plant sources
Patent: WO 011969-A 5 22-MAR-2001;
Large Scale Biology Corporation (US)
Location/Qualifiers
1..6446
/organism="Nicotiana benthamiana"
/db_xref="taxon:4100"

FEATURES
source

BASE COUNT 1873 a 1234 c 1563 g 1776 t

ORIGIN

alignment_scores:
Quality: 73.00 Length: 9
Ratio: 8.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x AX098418 ..

Align seg 1/1 to: AX098418 from: 1 to: 6446

4 LysTrpProTrpTrpProTrpArgArg 12
|||||
6213 AAGTGGCTTGTTGGTCATGCGCGCA 6239

seq_name: gb_htg:AP003754

seq_documentation_block:
LOCUS AP003754 129052 bp DNA HTG 14-JUN-2001
DEFINITION Oryza sativa chromosome 7 clone OJ1341_A08, *** SEQUENCING IN
PROGRESS ***, in ordered pieces.
ACCESSION AP003754
VERSION AP003754.1 GI:14422472
KEYWORDS HTG; HTGS; PHASE2.
SOURCE Oryza sativa
ORGANISM Oryza sativa
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Eriaristidoideae; Oryzaceae; Oryza.
1 (sites)
Sasaki,T., Matsumoto,T. and Yamamoto,K.
Oryza sativa nipponbare(GA3) genomic DNA, chromosome 7, BAC
clone:OJ1341_A08
Published Only in DataBase (2001) In press
2 (bases 1 to 129052)
Sasaki,T., Matsumoto,T. and Yamamoto,K.
Direct Submission
Submitted (13-JUN-2001) Takuji Sasaki, National Institute of
Agrobiological Resources, Rice Genome Research Program, Kannondai
2-1-2, Tsukuba, Ibaraki 305-8602, Japan
(E-mail:tsasaki@abrr.affrc.go.jp; URL:http://rgrp.dna.affrc.go.jp/;
Tel:81-298-38-7441, Fax:81-298-38-7468)
The nucleotide sequence of this BAC clone was generated by
combining Monsanto and RGP-Japan sequencing data.
NOTE: It currently consists of 1 contigs. Gaps between the contigs
are represented as runs of N. The order of the pieces is believed
to be correct as given, however the sizes of the gaps between them
are based on estimates that have provided by the submitter. This
sequence will be replaced by the finished sequence as soon as it is
available and the accession number will be preserved.
* NOTE: This is a 'working draft' sequence.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
Location/Qualifiers
1..129052

FEATURES
source

/organism="Oryza sativa"
/cultivar="Nipponbare"
/db_xref="taxon:4530"
/chromosome="7"
/clone="OJ1341_A08"

BASE COUNT 35088 a 28842 c 28921 g 36151 t 50 others

ORIGIN

alignment_scores:
Quality: 68.00 Length: 11
Ratio: 6.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:
US-09-444-281-35 x AP003754/rev ..

Align seg 1/1 to reverse of: AP003754 from: 1 to: 129052

3 LysLysTrpProTrpTrpProTrpArgArgLys 13
:::|||||
53246 CCGCGCTGGCTTGTTGGCTTGCGCGCGCGCG 53214

seq_name: gb_ro:MM008210

seq_documentation_block:
LOCUS MM008210 2651 bp mRNA ROD 31-OCT-1995
DEFINITION Mus musculus tropoelastin mRNA, complete cds.
ACCESSION U08210
VERSION U08210.1 GI:473273
KEYWORDS
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 2651)
Wyder,K.S., Sechler,J.L., Boyd,C.D. and Passmore,H.C.
Use of an intron polymorphism to localize the tropoelastin gene to
mouse chromosome 5 in a region of linkage conservation with human
chromosome 7
Genomics 23 (1), 125-131 (1994)
95130069
2 (bases 1 to 2651)
REFERENCE MEDLINE
AUTHORS Boyd,C.D.
TITLE Direct Submission
JOURNAL Submitted (30-MAR-1994) Charles D. Boyd, Department of Surgery,
UMDNJ - Robert Wood Johnson Medical School, 51 French St., New
Brunswick, NJ 08903, USA

FEATURES
source
Location/Qualifiers
1..2651
/organism="Mus musculus"
/strain="BALB/c"
/db_xref="taxon:10090"
/chromosome="5"
/tissue_type="lung"
/dev_stage="adult"
1..2583
/codon_start=1
/product="tropoelastin"
/protein_id="AAA80155.1"
/db_xref="GI:473274"
/translation="MAGLTVAVPQEVLLILLNLHPAOPGVGAVPGGLPGVPG
GVYYPGAGIGLGGGGALGPGRPKPPDAGLLTFFGAPGGLGAGPAGLGA
TPFGAGALVPGGAGAAAYKAAKAAAGAGCGVGPVGVGAVPGVGVGVG
GVGAGVGVGIGIGIGAGVAVVPGAGIGAGRGKRGVGVGLPVPVGVLP
TGARPVGAVLVGVPGVGVKAKAVGGGAGFSGIFGVPGGQDPGLGPIYIKAPL
GGIGLPTNGLPYGVAGAGKAGTPTGTGVSAAAAAATAAAYGAGAGVLPAGI
GCTPGAGALVGTGIGIAGATPAAAAAATAAATAAGAGLVPGGCVLPAGI
PGVGGIPGVGIGIPVGPDIIGDPGVGPGAVSPPAAATAAATAKARYGAGVGLPT
GVGAGFPGVGVGAGAGIGAGSPAAAAAATAAATAKARYGAGAGALGLVPGAVGLP
VAVAVGAGVPGVAGTPAAAAAATAAATAAATAKARYGAGVGVGAGVGVGAG
PGVGVGAVTGTGAGTGGAGGAGSPAAKASAKAAKAAKARYRAAGVAGVPGVAG
VPGFAGVAVPGFAGAGVPGFAGAGVPGFAGAVPGSLAAKAAKARYAAGAGLGGPG

BASE COUNT 410 a 623 c 1029 g 589 t
ORIGIN
alignment_scores:
Quality: 66.00 Length: 10
Ratio: 7.333 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000
alignment_block:
US-09-444-281-35 x MMU08210
Align seg 1/1 to: MMU08210 from: 1 to: 2651
4 LysTrpProTTrpProTArgLys 13
:::|||||
2025 AGTGGCCTTGCTGGCCCTGGAGCTCGG 2054
seq_name: gb_ro:AF289665
seq_documentation_block:
LOCUS AF289665 107257 bp. DNA ROD 14-AUG-2000
DEFINITION Mus musculus EIF4H gene, partial cds; LIMK1 gene, complete cds; and
ELN gene, partial cds.
ACCESSION AF289665
VERSION AF289665.1 GI:9800517
KEYWORDS house mouse.
SOURCE Mus musculus
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 107257)
AUTHORS Green,E.D.
TITLE Direct Submission
JOURNAL Submitted (26-JUL-2000) Genome Technology Branch, National Human
Genome Research Institute, 49 Convent Dr. Rm. 2A02, Bethesda, MD
20892, USA
FEATURES
source
1..107257
Location/Qualifiers
/organism="Mus musculus"
/strain="129/Sv"
/db_xref="taxon:10090"
/chromosome="5"
/clone="42120"
/clone_lib="Research Genetics CIB-BAC library"
/product="EIF4H"
/protein_id="AAFG9336.1"
/db_xref="GI:9800519"
/translation="MADFDYDDRAVSFGGRG"
/complement(join(24480..25812,25920..26077,26336..26391,
28274..28430,30220..30285,30530..30589,32612..32743,
33634..33720,34257..34440,37341..37507,37617..37722,
38745..38951,41098..41207,41439..41577,55071..55167,
56768..57026))
/product="LIMK1"
/complement(join(25650..25812,25920..26077,26336..26391,
28274..28430,30220..30285,30530..30589,32612..32743,
33634..33720,34257..34440,37341..37507,37617..37722,
38745..38951,41098..41207,41439..41577,55071..55167,
56768..57026))
/codon_start=1
/evidence=not_experimental
/product="LIMK1"
/protein_id="AAFG9334.1"
/db_xref="GI:9800518"

/translation="MRLTLCTWRRERMGESESLPVCASCGORLYDQVLOALNAD
WHADECRCECSVSLSHOYERKDGOLFCKKDWYAGVSGCHGSEHITLVLVWGEFL
KYHRECFICLGNFICDDDTYTHSHSLSCGCGYVYVYVETIOLIPDSRGLP
HVTYLVISFASAHKRGKLSVSDIPPGPFGCGTBSHTYTRVQGVDPGCSPPDKNTH
VGRILTEINGTPIRNVPLDEIDLIOETSRLLQTLLEHPHDSLGHPVSDPSLP
VHTPSQQAASSAKOKPVLRSCLSDTSPGTSLSLSPASQSRDLRSLSLRYVCPHPIE
RPSDLHGEVLGKGFQGAIKVTHRETFGWMMKELIRFEDEFORFLEVKMRCLE
HNVLFKIFGLVKDKRLNFITEYIKGTLGIIKNMSOYPMGSRVFAKDELSMAY
LHSMNIHHRDLSHNCILVRENRYVADDFGLARIMIDERNOSDLSTLKKPKRKYT
VYGNPYWMAPEMNGRSYDEKVDSEFGLICEITIRVANDPYLERTMDPGLVNCF
LDRYCPNCPPEFFPIITVRCDDIDPEKRSFVKLEQWLETLRHLHSLPLGQDLQ
ERGFETRYRGESESLPAHPEVPD"
complement(join(72144..72255,72961..73011,73719..73748,
74172..74216,74796..74846,74997..75035,75941..76081,
76252..76314,76403..76504,78538..78576,79493..79651,
80837..80896,80981..81121,82428..82485,83678..83779,
84115..84162,84859..84948,85423..85476,86452..86598,
87420..87479,87564..87653,90048..90095,90456..90510,
90895..90936,93350..93421,94200..94229,94899..94970,
96821..96868,97559..97621,97702..97863,98076..98177,
99496..99531,100964..100993,104734..104766,
105595..>105630))
/note="GenScan: P1=0.902, P2=0.979, P3=0.899, P4=0.797,
P5=0.640, P6=0.421, P7=0.897, P8=0.912, P9=0.991,
P10=0.999, P11=0.630, P12=0.813, P13=0.935, P14=0.767,
P15=0.773, P16=0.567, P17=0.806, P18=0.998, P19=0.999,
P20=0.966, P21=0.978, P22=0.980, P23=0.999, P24=0.966,
P25=0.579, P26=0.757, P27=0.661, P28=0.985, P29=0.952,
P30=0.228, P31=0.203, P32=0.451, P33=0.463, P34=0.528,
P35=0.228, P36=0.969, P37=0.995, P38=0.905, P39=0.655,
P40=0.986, P41=0.973, P42=0.930, P43=0.967, P44=0.995,
P45=0.999, P46=0.351, P47=0.994, P48=0.534, P49=0.787,
P50=0.943, P51=0.94"
/product="ELN"
complement(join(72212..72255,72961..73011,73719..73748,
74172..74216,74796..74846,74997..75035,75941..76081,
76252..76314,76403..76504,78538..78576,79493..79651,
80837..80896,80981..81121,82428..82485,83678..83779,
84115..84162,84859..84948,85423..85476,86452..86598,
87420..87479,87564..87653,90048..90095,90456..90510,
90895..90936,93350..93421,94200..94229,94899..94970,
96821..96868,97559..97621,97702..97863,98076..98177,
99496..99531,100964..100993,104734..104766,
105595..>105630))
/codon_start=3
/product="ELN"
/protein_id="AAFG9336.1"
/db_xref="GI:9800520"
/translation="AGTIGLGGGALGPGKPKPGAGLITFGAGPGGLGAGPGA
GIGAPFAGTFFPGAGALVPGGAGAAVYAKAAKAGGLGAGVPGVPGVPGVPGV
VGGVPGVPGVPGVPGVPGVPGVPGVPGVPGVPGVPGVPGVPGVPGVPGVPGV
YFGVLPFGTGAFFPGVGLPEVPTGTGVAKAPGGGAGRIGIPGVFGQDQVPLG
YPLKAPKLPGGYGLPYTNGLRPGVAGAGKAGKAPGTGVSQAAAKAAKAYGAG
ACVLPDGVGGGIPGAGAIPIGLIGAGCTPAAAKAAKAAKAAKAAKAYGAG
VRLPGAGIPGVGIPGVGIPGVGIPGVGIPGVGIPGVGIPGVGIPGVGIPGVG
GGVGLPTGYGAGGPPGYGVGAGAGLGGSPAAAAAAKAAKAAKAAKAAKAYGAG
VPGALPGAVPAVAGVPGAGTGPAAAPAAAAAAKAAKAAKAAKAAKAYGAG
GGV
PGGAGAGVPGFAGAGVPGFAGAGVPGFAGAGVPGFAGAGVPGFAGAGVPGFAGAG
AGGLGAGLGGAGLGGAGLGGAGLGGAGLGGAGLGGAGLGGAGLGGAGLGGAGLGGAG
GLGGVYARPPPGGVVAAARPGFLSPIYPGGAGLGGVGGKPKPKPGALGALGVGG
GGGKSGGKKR"

BASE COUNT 26300 a 26393 c 26598 g 27966 t
ORIGIN

alignment_scores:
Quality: 66.00 Length: 10
Ratio: 7.333 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000
alignment_block:

US-09-444-281-35 x AF289665/rev
Align seg 1/1 to reverse of: AF289665 from: 1 to: 107257

4 LysTrpProTrpTrpProTrpArgArgLys 13
:::|||||

76497 AGGTGGCTTGTCGTCGAGGTCGCG 76468

seq_name: gb_htg:AC091250

seq_documentation_block:

LOCUS AC091250 200849 bp DNA HTG 11-APR-2001
DEFINITION Mus musculus chromosome 5 clone RP23-315E2 strain C57BL6/J, WORKING
DRAFT SEQUENCE, 7 unordered pieces.

AC091250
AC091250.1 GI:13592171

HTG: HTGS_PHASE1; HTGS_DRAFT.
house mouse.

SOURCE

ORGANISM

Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 200849)
Aylee, K., Beckstrom-Sternberg, S.M., Benjamin, B., Blakesley, R.W.,
Boutard, G.G., Brinkley, C., Brooks, S., Dietrich, N.L., Granite, S.,
Guan, X., Gupta, J., Ho, S.-L., Idol, J.R., Karlins, E., Lee-Lin, S.-O.,
Legaspi, R., Lim, M., Maduro, Q.L., Maduro, V.B., Mastello, C.,
Mastrian, S.D., McCloskey, J.C., McPowell, J., Pearson, R., Prasad, A.,
Shevchenko, Y., Snyder, B., Stantrop, S., Thomas, J.W., Thomas, P.J.,
Thompson, E.E., Touchman, J.W., Tsurguev, C., Vogt, J.L., Walker, M.A.,
Wetherby, R.D., Zhang, L.H. and Green, E.D.

NIH Comparative Sequencing Initiative
Unpublished

2 (bases 1 to 200849)
Green, E.D.

Direct Submission
Submitted (11-APR-2001) NIH Intramural Sequencing Center, 8717
Grovermont Circle, Gaithersburg, MD 20877, USA

Center: NIH Intramural Sequencing Center
Genome Center

Center code: NISC
Web site: http://www.nisc.nih.gov

Contact: nisc.mouse@nhgri.nih.gov
Project Information

Center project name: aty
Center clone name: 315E02
Summary Statistics
Sequencing vector: plasmid; n/a; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap, version 0.990319
Consensus quality: 197727 bases at least Q40
Consensus quality: 198423 bases at least Q30
Consensus quality: 198823 bases at least Q20
Insert size: 202000; agarose-fp
Insert size: 200249; sum-of-contigs
Quality coverage: 9.87x in Q20 bases; sum-of-contigs
Quality coverage: 9.95x in Q20 bases; sum-of-contigs

NOTE: This is a 'working draft' sequence. It currently
consists of 7 contigs. The true order of the pieces
is not known and their order in this sequence record is
arbitrary. Gaps between the contigs are represented as
runs of N, but the exact sizes of the gaps are unknown.
This record will be updated with the finished sequence
as soon as it is available and the accession number will
be preserved.

1 2812: contig of 2812 bp in length
* 2813 2812: gap of unknown length
* 2913 9301: contig of 6389 bp in length
* 9302 9401: gap of unknown length
* 9402 17173: contig of 7772 bp in length
* 17174 17273: gap of unknown length
* 17274 26044: contig of 8771 bp in length
* 26045 26144: gap of unknown length

FEATURES
source
1. 200849
/organism="Mus musculus"
/strain="C57BL6/J"
/db_xref="taxon:10090"
/chromosome="5"
/clone="RP23-315E2"
/clone_1ib="RPC1 mouse BAC library 23"
1. 2812
/note="assembly_fragment"
2913 9301
/note="assembly_fragment"
9402 17173
/note="assembly_fragment"
17274 26044
/note="assembly_fragment"
26145 65929
/note="assembly_fragment"
clone_end:SP6
vector_side:right"
66030 109036
/note="assembly_fragment"
109137 200849
/note="assembly_fragment"
clone_end:SP6
vector_side:right"

BASE COUNT 53444 a 49164 c 49019 g 48614 t 608 others
ORIGIN
alignment_scores:
Quality: 66.00 Length: 10
Ratio: 7.333 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000
alignment_block:
US-09-444-281-35 x AC091250 ..
Align seg 1/1 to: AC091250 from: 1 to: 200849

4 LysTrpProTrpTrpProTrpArgArgLys 13
:::|||||
148453 AGGTGGCTTGTCGTCGAGGTCGCG 148482

seq_name: gb_htg:AC020365

seq_documentation_block:

LOCUS AC020365 27780 bp DNA HTG 03-JAN-2000
DEFINITION Drosophila melanogaster, *** SEQUENCING IN PROGRESS ***
pieces.

AC020365
AC020365.1 GI:6664532

HTG: HTGS_PHASE2.
fruit fly

ORGANISM
Drosophila melanogaster

Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

1 (bases 1 to 27780)
Adams, M. and Venter, J.C.

Direct Submission
Submitted (30-DEC-1999) Celera Genomics, 45 West Gude Drive,
Rockville, MD, USA

This sequence was identified as CDL:10212917 by the submitter.
For more information on this record e-mail to fly@celera.com.

NOTE: This is a 'working draft' sequence.
This sequence will be replaced
by the finished sequence as soon as it is available and

* the accession number will be preserved.

FEATURES
source 1. 27780
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"

BASE COUNT 7869 a 6128 c 6190 g 7593 t
ORIGIN

alignment_scores:
Quality: 65.00 Length: 15
Ratio: 5.909 Gaps: 1
Percent Similarity: 73.333 Percent Identity: 60.000

alignment_block:
US-09-444-281-35 x AC020365 ..

Align seg 1/1 to: AC020365 from: 1 to: 27780

3 LysLysTTPPro.....TTPTrProTTPArGArGlys 13
||||:|||||
10280 AAGAGGTGGCCCTGGAGAAAGGTGGTGGCCCTGGAGAGAGAA 10324

seq_name: gb_in:AC008316

seq_documentation_block:

LOCUS AC008316 160817 bp DNA INV 06-MAR-2001
DEFINITION Drosophila melanogaster, chromosome 3R, region 85F-85F, BAC clone
BACR23004, complete sequence.

ACCESSION AC008316
VERSION AC008316 GI:13236627
KEYWORDS HTG.

SOURCE
ORGANISM

Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 160817).

REFERENCE
AUTHORS

Celniker, S.E., Adams, M.D., Krommler, B., Tyler, D., Wan, K.H.,
Holt, R.A., Evans, C.A., Gocayne, J.D., Amanatides, P.G., Brandon, R.C.,
Rogers, Y., An, H., Baldwin, D., Banzon, J., Beeson, K.Y., Busam, D.A.,
Carlson, J.W., Center, A., Chape, M., Davenport, L.B., Dietz, S.M.,
Dodson, K., Dorselt, V., Doup, L.E., Doyle, C., Dresnek, D., Farfan, D.,
Fierlier, S., Frise, E., Galle, R.F., Gary, N.S., George, R.A.,
Gonzalez, M., Houch, J., Hoskins, R.A., Hostin, D., Howland, T.J.,
Ibegam, C., Jatali, M., Kruse, D., Li, P., Mattei, B., Moshirefi, A.,
McIntosh, T.C., Moy, M., Murphy, B., Nelson, C., Nelson, K.A., Nunoo, J.,
Paciel, J., Paragas, V., Park, S., Patel, S., Pfeiffer, B.,
Phouanavong, S., Pittman, G.S., Puri, V., Richards, S., Scheeler, F.,
Stapleton, M., Strong, R., Svirskaas, R., Tector, C., Williams, S.M.,
Zaveri, J.S., Smith, H.O., Rubin, G.M. and Venter, J.C.
Sequencing of Drosophila chromosome 3R, region 85F-85F
Unpublished
2 (bases 1 to 160817)

TITLE
JOURNAL
REFERENCE
AUTHORS

Celniker, S.E., Agapayni, A., Arcaina, T.T., Baxter, E., Blazek, R.G.,
Butenoff, C., Chape, M., Chavez, C., Chew, M., Ciesiolka, L.,
Doyle, C.M., Farfan, D.E., Galle, R., George, R.A., Harris, N.L.,
Hoskins, R.A., Houston, K.A., Hummasti, S.R., Karia, K., Kearney, L.,
Kim, E., Lee, B., Lewis, S., Li, P., Lomocan, M.A., Mazda, P.,
Moshirefi, A.R., Moshirefi, M., Nixon, K., Paciel, J.M., Park, S.,
Pfeiffer, B., Poon, L., Sequeira, A., Sethi, H., Snir, E.,
Svirskaas, R.R., Wan, K.H., Weinburg, T., Zhang, R., Zieran, L.L. and
Rubin, G.M.

TITLE
JOURNAL

Submitted (02-AUG-1999) Drosophila Genome Center, Lawrence Berkeley
Laboratory, MS 64-121, Berkeley, CA 94720, USA
On Mar 6, 2001 this sequence version replaced gi:6957904.

COMMENT

Sequence submitted by:
Berkeley Drosophila Genome Project
Lawrence Berkeley National Laboratory, MS 64-121
Berkeley, CA 94720
This sequence was assembled using end sequences from a whole genome
shotgun and from subclones of this BAC and its neighboring clones.

For further information about this sequence, including its location
and relationship to other sequences, please visit our sequence
archive Web site (<http://www.fruitfly.org/sequence/>) or send email
to bdg@fruitfly.berkeley.edu.

FEATURES
source 1. 160817
/organism="Drosophila melanogaster"
/strain="y: cn bw sp"
/db_xref="taxon:7227"
/chromosome="3R"
/map="85F-85F"
/clone.lib="RPCI-98 (Roswell Park Cancer Institute
Drosophila melanogaster BAC library, partial EcoRI in
pBAC3.6)"

BASE COUNT 4532 a 35217 c 34880 g 45188 t
ORIGIN

alignment_scores:
Quality: 65.00 Length: 15
Ratio: 5.909 Gaps: 1
Percent Similarity: 73.333 Percent Identity: 60.000

alignment_block:
US-09-444-281-35 x AC008316 ..

Align seg 1/1 to: AC008316 from: 1 to: 160817

3 LysLysTTPPro.....TTPTrProTTPArGArGlys 13
||||:|||||
18897 AAGAGGTGGCCCTGGAGAAAGGTGGTGGCCCTGGAGAGAGAA 18941

seq_name: gb_in:AC008315

seq_documentation_block:

LOCUS AC008315 177028 bp DNA INV 17-FEB-2001
DEFINITION Drosophila melanogaster, chromosome 3R, region 85E-85E, BAC clone
BACR30F01, complete sequence.

ACCESSION AC008315
VERSION AC008315 GI:12957660
KEYWORDS HTG.

SOURCE

ORGANISM

Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 177028)

REFERENCE
AUTHORS

Celniker, S.E., Adams, M.D., Krommler, B., Tyler, D., Wan, K.H.,
Holt, R.A., Evans, C.A., Gocayne, J.D., Amanatides, P.G., Brandon, R.C.,
Rogers, Y., An, H., Baldwin, D., Banzon, J., Beeson, K.Y., Busam, D.A.,
Carlson, J.W., Center, A., Chape, M., Davenport, L.B., Dietz, S.M.,
Dodson, K., Dorselt, V., Doup, L.E., Doyle, C., Dresnek, D., Farfan, D.,
Fierlier, S., Frise, E., Galle, R.F., Gary, N.S., George, R.A.,
Gonzalez, M., Houch, J., Hoskins, R.A., Hostin, D., Howland, T.J.,
Ibegam, C., Jatali, M., Kruse, D., Li, P., Mattei, B., Moshirefi, A.,
McIntosh, T.C., Moy, M., Murphy, B., Nelson, C., Nelson, K.A., Nunoo, J.,
Paciel, J., Paragas, V., Park, S., Patel, S., Pfeiffer, B.,
Phouanavong, S., Pittman, G.S., Puri, V., Richards, S., Scheeler, F.,
Stapleton, M., Strong, R., Svirskaas, R., Tector, C., Williams, S.M.,
Zaveri, J.S., Smith, H.O., Rubin, G.M. and Venter, J.C.
Sequencing of Drosophila chromosome 3R, region 85E-85E
Unpublished
2 (bases 1 to 177028)

TITLE
JOURNAL
REFERENCE
AUTHORS

Celniker, S.E., Agapayni, A., Arcaina, T.T., Baxter, E., Blazek, R.G.,
Butenoff, C., Chape, M., Chavez, C., Chew, M., Ciesiolka, L.,
Doyle, C.M., Farfan, D.E., Galle, R., George, R.A., Harris, N.L.,
Hoskins, R.A., Houston, K.A., Hummasti, S.R., Karia, K., Kearney, L.,
Kim, E., Lee, B., Lewis, S., Li, P., Lomocan, M.A., Mazda, P.,
Moshirefi, A.R., Moshirefi, M., Nixon, K., Paciel, J.M., Park, S.,
Pfeiffer, B., Poon, L., Sequeira, A., Sethi, H., Snir, E.,
Svirskaas, R.R., Wan, K.H., Weinburg, T., Zhang, R., Zieran, L.L. and
Rubin, G.M.

TITLE Direct Submission
JOURNAL Submitted (02-AUG-1999) Drosophila Genome Center, Lawrence Berkeley Laboratory, MS 64-121, Berkeley, CA 94720, USA
COMMENT On Feb 17, 2001 this sequence version replaced gi:7248925.
Sequence submitted by:
Berkeley Drosophila Genome Project
Lawrence Berkeley National Laboratory, MS 64-121
Berkeley, CA 94720
This sequence was assembled using end sequences from a whole genome shotgun and from subclones of this BAC and its neighboring clones. For further information about this sequence, including its location and relationship to other sequences, please visit our sequence archive Web site (<http://www fruitfly.org/sequence/>) or send email to bdbg@fruitfly.berkeley.edu.
FEATURES
Location/Qualifiers
Source
1..177028
/organism="Drosophila melanogaster"
/strain="y; cn bw sp"
/db_xref="taxon:7227"
/chromosome="3R"
/map="85E-85E"
/clone_lib="BACRCF01 (D3836)"
/clone_id="RPCL1-98 (Roswell Park Cancer Institute Drosophila melanogaster BAC library, partial EcoRI in pBac3.6)"
BASE COUNT 48567 a 39865 c 40667 g 47929 t
ORIGIN

alignment_scores:
Quality: 65.00 Length: 15
Ratio: 5.909 Gaps: 1
Percent Similarity: 73.33 Percent Identity: 60.000

alignment_block:
US-09-444-281-35 x AC008315 ..
Align seg 1/1 to: AC008315 from: 1 to: 177028

3 lyslystTPPro.....TrrTPProTrPARGATGLys 13
|||||:||||||| |||||:|||||:|||||
126050 AAGAGGTGGCCCTGGAGCGAAGAAGTGTCGCCCTGGAGAGAA 126094

seq_name: gb_in:AEO03684 ?
seq_documentation_block:
LOCUS AEO03684 219579 bp DNA INV 05-OCT-2000
DEFINITION Drosophila melanogaster, genomic scaffold 142000013386035 section 9 of 105, complete sequence.
ACCESSION AEO03684 AEO02708
VERSION AE003684.2 GI:10726416
KEYWORDS HTG.
SOURCE fruit fly.
ORGANISM Drosophila melanogaster
Eukaryota; Neoptera; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Metazoa; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephyridioidea; Drosophilidae; Drosophila.
1 (bases 1 to 219579).
Adams,M.D., Celniker,S.E., Holt,R.A., Evans,C.A., Gocayne,J.D.,
Amantides,P.G., Scherer,S.E., Li,P.W., Hoskins,R.A., Galie,R.F.,
George,R.A., Lewis,S.E., Richards,S., Ashburner,M., Henderson,S.N.,
Sutton,G.G., Wootman,J.R., Yandell,M.D., Zhang,Q., Chen,L.X.,
Brandon,R.C., Rogers,Y.H., Blazer,R.G., Chample,M., Pfeiffer,B.D.,
Wan,K.H., Doyle,C., Baxter,E.G., Helt,G., Nelson,C.R., Gaber
Miklos,G.L., Abriil,J.F., Agbayani,A., An,H.J.,
Andrews-Pfanczuch,C., Baldwin,D., Ballev,R.M., Basu,A.,
Baxendale,J., Bayraktaroglu,L., Beasley,E.M., Beeson,K.Y.,
Benos,P.V., Bergman,B.P., Bhandari,D., Bolshakov,S., Borokova,D.,
Botchan,M.R., Bouck,J., Brokstein,P., Brotlier,P., Buttis,K.C.,
Busam,D.A., Butler,H., Cadieu,E., Center,A., Chandra,I.,
Cherry,J.M., Cawley,S., Dahlke,C., Davenport,L.B., Davies,P., de
Pablos,B., Delcher,A., Deng,Z., Mays,A.D., Dew,I., Dietz,S.M.,
Dodson,K., Doup,L.E., Downes,M., Dugan-Rocha,S., Dunkov,B.C.,

FEATURES	COMMENT
source	location/Qualifiers
1. 219579	
/organism="Drosophila melanogaster"	
/db_xref="taxon:7227"	
/chromosome="3R"	
/complement(join(608..879,945..1093,1160..1371,1428..1672,1738..2464,2521..2728,2789..3377,3441..3601,4534..5054))	
/gene="CG9448"	
/product="CT26774"	
/db_xref="FLYBASE:Fban0009448"	
/db_xref="FLYBASE:FBgn0037734"	
complement(<608..5054)	
/gene="CG9448"	
/note="CG9448 gene product"	
/codon_start=1	
/db_xref="FLYBASE:Fban0009448"	
/db_xref="FLYBASE:FBgn0037734"	
/protein_id="AF54429.1"	
/db_xref="GI:7299233"	
/translation="MCDTKDDAQKWCETCTYNYPSISKCTMKCQASKPLLNIDFRLSPQESCVAEEAAVEAVMSPSPSTCSLQPOSARQSNVADSEKMKCVTYLNWPSLSCVOCCTKRGGEALERKMDNEDNGRAGSLADLITSGEENLAKPQOLIGTASHRLSLSRGIDDAVHLNNLANASHQSOHORPQLOOMOLOLOPOOPRSSSAVAPOOQKOCYKQACNOCSTYENNPWPSIKCSCCKTREFEISGSQNDLAASSLNSQEEQOOLQOPNVTVSYVNNSEFNKKHLYQDLSSETTINNCDTLQERQERQIRQVQOMNACLGVENNYSAEVALSCGNGPARELSTETLAINCLRNARSAFGRHLILAIRHREMLPMLDIQISSGPGIKRVSVAADPLADIRPFPNNTLRKSGPCHYVOKHAFPAEIEIPPIQEOPLDYDLRLDQAKQOLETPPANTLSLETISARSRFVLMNRASGQCLDLSAQATAGVDRNDINLRALADLHLCQGHVFRKWEYEMLOASLHFTLDSQOEEDMSTLLSLAQOGSSLDLHFAHLHILRLPIIYGVKYVSPGCDIGVAFEGEVYLPFWNDQCNKSPALATYTRGHFSALVPMPEFTTIDGRDVEYTYLIMQCEKLKLPIHLTQSEVNGENESMMRWMLDVCVTGGGLLVAAQCKUSKRLPVLAAQMLEEMLNHRTRIAQVITAFIRPQTLTHSDSDSE"	
/gene="dmt"	
join(5193..5743,5803..7634,7920..8728)	
CDS	
gene	
mRNA	
REFERENCE	
AUTHORS	
JOURNAL	
TITLE	
MEDLINE	
JOURNAL	
2 (bases 1 to 219579)	
Adams,M.D., Celisner,S.E., Gibbs,R.A., Rubin,G.M. and Venter,C.J.	
Direct Submission	
Submitted (21-MAR-2000) Celera Genomics, 45 West Gude Drive, Rockville, MD, USA	
On Oct 9, 2000 this sequence version replaced gi:7299232.	
Location/Qualifiers	
1. 219579	
/organism="Drosophila melanogaster"	
/db_xref="taxon:7227"	
/chromosome="3R"	
/complement(join(608..879,945..1093,1160..1371,1428..1672,1738..2464,2521..2728,2789..3377,3441..3601,4534..5054))	
/gene="CG9448"	
/product="CT26774"	
/db_xref="FLYBASE:Fban0009448"	
/db_xref="FLYBASE:FBgn0037734"	
complement(<608..5054)	
/gene="CG9448"	
/map="85E5-85E6"	
/db_xref="FLYBASE:Fban0009448"	
/db_xref="FLYBASE:FBgn0037734"	
complement(join(801..879,945..1093,1160..1371,1428..1672,1738..2464,2521..2728,2789..3377,3441..3568))	
/gene="CG9448"	
/note="CG9448 gene product"	
/codon_start=1	
/db_xref="FLYBASE:Fban0009448"	
/db_xref="FLYBASE:FBgn0037734"	
/protein_id="AF54429.1"	
/db_xref="GI:7299233"	
/translation="MCDTKDDAQKWCETCTYNYPSISKCTMKCQASKPLLNIDFRLSPQESCVAEEAAVEAVMSPSPSTCSLQPOSARQSNVADSEKMKCVTYLNWPSLSCVOCCTKRGGEALERKMDNEDNGRAGSLADLITSGEENLAKPQOLIGTASHRLSLSRGIDDAVHLNNLANASHQSOHORPQLOOMOLOLOPOOPRSSSAVAPOOQKOCYKQACNOCSTYENNPWPSIKCSCCKTREFEISGSQNDLAASSLNSQEEQOOLQOPNVTVSYVNNSEFNKKHLYQDLSSETTINNCDTLQERQERQIRQVQOMNACLGVENNYSAEVALSCGNGPARELSTETLAINCLRNARSAFGRHLILAIRHREMLPMLDIQISSGPGIKRVSVAADPLADIRPFPNNTLRKSGPCHYVOKHAFPAEIEIPPIQEOPLDYDLRLDQAKQOLETPPANTLSLETISARSRFVLMNRASGQCLDLSAQATAGVDRNDINLRALADLHLCQGHVFRKWEYEMLOASLHFTLDSQOEEDMSTLLSLAQOGSSLDLHFAHLHILRLPIIYGVKYVSPGCDIGVAFEGEVYLPFWNDQCNKSPALATYTRGHFSALVPMPEFTTIDGRDVEYTYLIMQCEKLKLPIHLTQSEVNGENESMMRWMLDVCVTGGGLLVAAQCKUSKRLPVLAAQMLEEMLNHRTRIAQVITAFIRPQTLTHSDSDSE"	
/gene="dmt"	
join(5193..5743,5803..7634,7920..8728)	

```

/note="Nucleotide sequence of the Celera sequence differs
from the published sequence for this transcript."
/db_xref="C24655"
/db_xref="FLYBASE:FBan0008374"
/db_xref="FLYBASE:FBgn0016792"
<193>..>8728
/gene="dmt"
/note="CG8374"
/map="85E5-85E6"
/db_xref="FLYBASE:FBan0008374"
/db_xref="FLYBASE:FBgn0016792"
join(5263..5743,5803..7634,7920..8180)
/gene="dmt"
/note="dmt gene product: Nucleotide sequence of the Celera
sequence differs from the published sequence for this
transcript"
/codon_start=1
/db_xref="FLYBASE:FBan0008374"
/db_xref="FLYBASE:FBgn0016792"
/protein_id="AAF54430.1"
/db_xref="GI:7299234"
/translation="MYRTPRVPCVSPDVNTATRRNPGKPKKOSIGADLSTTISKRCR
PKLSTIGADLTTIRKPRPKLGADLTTIRKPRPKLSTKOSITALEPERRIKKC
VYKIRIKKNGVSEISRNDAVDASTFEPARNTMYTPSKVOSTQPONIKENA
GEBTSCQORIRKSPSGPAEISPKPRPFHQDPLTRTSSVTRVPELDSOI
EDSDREDAADILORLVKDGACVWRKTIKPRAKKRVGNRRKYSTKD
EEPRVYVYKSTIEPRKRSSTIEPEEDYSNDEHGYQVLTIEPVQVYVEPST
KAHBGAYSNLASVHNOTQONPLSTDRRELINAKQLSTPLNRRAPVIDVS
ATTALSLPIAHSPPNAVRTAGKSPWRVSDSDPLPTMFGFNNSQSPSDHVR
HYVVPSPVEHENIPHEESICPPLHEQNDNSANDNEPPTISKMSINDOSR
ENAEKRVHLNPNRRTRKRPDKDINILEVTLPSMKKNVATVKEITPTVRAPR
MATSSPAORSOTRSNLFQDDPACEDIDIKSTTPSKGIAPTSMISNREVTALARN
OREVYVNGEELIPCEINPEKENTNEPRTSSQNIQGEDEFTTESQDSPPANTGSO
NWNLDKRLRLAELPRDCELPQVSTYSTRDCLGSHKQDIRIMLSTMAPRP
ARRKPAAPARSMGLFRVODEPEQSFQDKQPRRYVERQQRKKRKOVLITYESE
SEDEEDQSDHSDLSPEKKRHHVRRRDIEHEAKLEQFVTSFNKCEVEKEPVLI
E"
complement(join(8675..9217,9277..9507,9574..9721,
9777..9983,10044..10369,10427..10507,10570..12329,
12384..12594,12684..13156,13217..13450,13530..13674,
13773..17273,17361..17551,17614..17852,17913..18130,
18190..18469,18533..18730,19517..19631))
/gene="hyd"
/note="Nucleotide sequence of the Celera sequence differs
from the published sequence for this transcript."
/product="C126854"
/db_xref="FLYBASE:FBan0009484"
/db_xref="FLYBASE:FBgn0002431"
complement(<8675..>19631)
/gene="hyd"
/note="CG9484"
/map="85E5-85E6"
/db_xref="FLYBASE:FBan0009484"
/db_xref="FLYBASE:FBgn0002431"
complement(join(11575..12329,12384..12594,12684..13156,
13217..13450,13530..13674,13773,17361..17551,
17614..17852,17913..18130,18190..18469,18533..18730,
19517..19569))
/gene="hyd"
/note="hyd gene product: Nucleotide sequence of the Celera
sequence differs from the published sequence for this
transcript"
/codon_start=1
/db_xref="FLYBASE:FBan0009484"
/db_xref="FLYBASE:FBgn0002431"
/protein_id="AAF54431.1"
/db_xref="GI:7299235"
/translation="MOFLVPLPGSDDOFIERIREVSDKVRNGYSGHRIPELQKIPV
KEVIGPAHIGVLEDKGAPRFSEISSEKLDITKDACSTSGSGTASAKAPSS
RPAARBARLLRATGRSNTSCGCSRSCTGVTIGGSTSPPLTVPATVYPELISQA
EYVLQKSRLLIRLQRTNLDVNTLAVNNLSDDEDEADTEGADNVPEDLITSLD
NGSGDNNSVITIDSDGLFSEELFYSNIRLIFRISERNANANANADNSQSTTR
SSSGTALNGNSGLSAQISVNDREAFSSRWDRQYGGPRRWISKDDYTWEKADSKKK

```

```

EPSPMLSPIWISPELOPMPWPKSSVREKTIKALYSEFIALSEGDLGYOMRWSAPPYKS
EPENYHYPRKTVLSINVERVELISANIRCSVTEPRVATPMMDPOLGYIGALTEKSCV
AFNERPISDSITIKIYVCSLITVYKTESNNTYMKVCLPFDDRRLEMDKFRKTKPKYV
AIDINGAIVIMKRCMPYSGSITGFPCSGNGVPRVGLNSVMTFDVCKMLKININTN
SGVDSQAAGNMLNANGITPDKLPKSTAMPSPGSRKNOQSFNSNKSJDRIDMPP
SPASSTCSDTGSLVSHKRTKRAATKEDSNAPOEGRRKDELMEYKDYFEDKYGPGK
VLKVDDEFAVPRPAINAAVAANAATSTSTNTASTSKECKEDMOOCRLIREDY
OIFRTMSTRGROKOPKTIYNGDAGAOITLTVASRGRIKVKYKIGKTHVSLY
NLYNCKOENCLPPTDCNSFTGISTPONTIIMACNDSCSGSSTIVLBDGALYPLAKD
CUGSTIKDPQWFDLPVKSITMSTSTSLPALSVNLSKSKVCMPTALLEDTOKLMPHLRC
DVKNSFALGRLEDEODADTALVERECGARIIFHACVIMCASPSSNKSDPPDSPG
VRRKSLVGLSNTIPTVSTSAVSIAGASASSNENSSFAATSSSAAGSASTSR
DNRTNLRDMNRLINSIDVSAOSQOSOMANNEDHAYIPWAPETPAASNASSOSVVS
DSIEDISKITPSSSSMLSNIGKSPPTFDIADRRHALITIOQMVSVALRPYL
CHMLSTRKDAQOGOTPRMLVSCRAYEKGITLLNTILMSQDQKLEAMIFPSPRADO
SPHATICTNDTCSFTWTGADHNONIFECRTGGLTSCCTCEAVCGKHGDKQR
TAPTAAYCDCEWCKCKALJAGNITKRFALLCKIVSTDLVTFNSKGESILLFLIOTV
GROIYBQRYRFSVRVNVSTATGATGNNSVINSKTSAAEIDNDMPDPLEPFA
RKALERLIDMNAVRSMISGAEKGVPPNPDASASNSNSSEGFNMEIOTQHGSTLIDK
FTHSLVTKCTSDHDLTLTLVIRELONASVNSRKEAEVVRPALLVPRGUVPRPPE
THSSMLNDSIDLFSDPLAPSVESPSQILYHDGNDQSANFINIQONTDYVAMETI
RQASEEVEYINREANSNHDDELLENQRENDQDESDNDEFTNDAETESDDMOS
NOEYORSVQAGATVGSENDIGVLFLEDESDSQAODESDSESDQSDDEFNFOQ
LBRRSVNSNARSDLAPQTMQAMIRSDTARRSVRYPTGNNMVIDPMALRSTVPAST
TVTTPTEIEPHMTATSNLARAFAVITIROISELITSLIVNLINDITSLKINDAIA
VOAFVYKRLKATWDMFTYMDGTEAGLKGALITNTTDPNHPHLPLNLSAQSSQTP

```

alignment_scores:

Quality:	Length:
65.00	15
Ratio: 5.909	Gaps: 1
Percent Similarity: 73.333	Percent Identity: 60.000

alignment_block:

US-09-444-281-35 x AE003684 ..

Align seg 1/1 to: AE003684 from: 1 to: 219579

3 LysylsTTPPro.....TTPTrProTTPaTgaTglys 13

38362 AAGAGGTGGCCCTGAGGAAAGGTGGCCCTGAGGAGGAA 38406

seq_name: gb_Jr:MMA2IXCOA

seq_documentation_block: ROD 08-APR-1994

LOCUS MMA2IXCOA 19479 bp DNA

DEFINITION M.musculus alpha2 (IX) collagen gene, complete CDS.

ACCESSION 222923

VERSION 222923.1 GI:311949

KEYWORDS alpha2 (IX) collagen.

SOURCE house mouse.

ORGANISM Mus musculus

Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.

1 (bases 1 to 19479)

Perala,M., Elina,K., Metsaranta,M., Rosati,R., de Crombrughe,B.

and Vuorio,E.

The exon structure of the mouse alpha 2(IX) collagen gene shows

unexpected divergence from the mouse alpha 2(IX) collagen gene

unpublished divergence from the mouse alpha 2(IX) collagen gene

unpublished divergence from the mouse alpha 2(IX) collagen gene

unpublished divergence from the mouse alpha 2(IX) collagen gene

exon	15284..15337	/gene="alpha2 (IX) collagen"	9
	//label=ex22		0
	/number=22		0
	/evidence=experimental		0
	15343..15396		0
exon	/gene="alpha2 (IX) collagen"		9
	//label=ex23		0
	/number=23		0
	/evidence=experimental		0
	15674..15745		0
exon	/gene="alpha2 (IX) collagen"		9
	//label=ex24		0
	/number=24		0
	/evidence=experimental		0
	15897..15932		0
exon	/gene="alpha2 (IX) collagen"		9
	//label=ex25		0
	/number=25		0
	/evidence=experimental		0
	16069..16113		0
exon	/gene="alpha2 (IX) collagen"		9
	//label=ex26		0
	/number=26		0
	/evidence=experimental		0
	16207..16239		0
exon			9
alignment_scores:	64.00	Length:	9
Quality:	8.000	Gaps:	0
Ratio:	88.889	Percent Identity:	77.778

Align seg 1/1 to reverse of: MMA2IXCOA from: 1 to: 19479

```

5 TrpProTrrPTrrProTrrPargArgLys 13
  |||||
4355 TGGCCCTGGTGGCCCTGAGAGACCCGG 4329

```

seq_name: gb_pr:AL162423

```
seq_documentation_block:
```

DEFINITION Human DNA se

ACCESSION ALL62423

KEYWORDS: HTG.

ORGANISM Homo sapiens

Mammalia; Eu

AUTHORS Garner, P.

JOURNAL Submitted (0

COMMENT On Mar 5, 20

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above. This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30): an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the

assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em', EMBL; Sw', SWISSPROT; Tr', TrEMBL; Wp', WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 9, constructed by the Sanger Centre Chromosome 9 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/chr9> RP11-456621 is from the library RP11-2 constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/bacpac/home.htm> VECTOR: pBAC3.6

IMPORTANT: This sequence is not the entire insert of clone RP11-456621 It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap. The true left end of clone RP11-456621 is at 1 in this sequence. The true left end of clone RP11-49882 is at 88324 in this sequence. The true right end of clone RP11-297L11 is at 34844 in this sequence.

INES	location/Qualifiers
source	1. .88323
	/organism="Homo sapiens"
	/db_xref="taxon:9606"
	/chromosome="9"
	/clone="RP11-456D21"
	/clone_1kb="RPC1-11.2"
repeat_L_region	362. .588
repeat_L_region	/note="HA1 repeat: matches 595. .829 of consensus"
repeat_L_region	693. .1009
repeat_L_region	/note="LM6C repeat: matches 2085. .2405 of consensus"
repeat_L_region	1168. .1731
repeat_L_region	/note="LM4 repeat: matches 2630. .3220 of consensus"
repeat_L_region	1775. .2047
repeat_L_region	/note="L2 repeat: matches 2248. .2526 of consensus"
repeat_L_region	2051. .2215
repeat_L_region	/note="MIR repeat: matches 82. .252 of consensus"
repeat_L_region	2443. .2788
repeat_L_region	/note="LM4 repeat: matches 5399. .5754 of consensus"
repeat_L_region	2889. .3184
repeat_L_region	2889. .3184
repeat_L_region	/note="A1y repeat: matches 1. .298 of consensus"
repeat_L_region	3306. .3446
repeat_L_region	/note="FLM_C repeat: matches 2. .133 of consensus"
repeat_L_region	3862. .4181
repeat_L_region	/note="A1u0 repeat: matches 2. .312 of consensus"
repeat_L_region	5285. .5366
repeat_L_region	/note="41 copies 2 mer aa 63* conserved"
repeat_L_region	5681. .5983
repeat_L_region	/note="A1u5 repeat: matches 1. .305 of consensus"
repeat_L_region	6211. .6508
repeat_L_region	/note="A1u0 repeat: matches 2. .298 of consensus"
repeat_L_region	6566. .6961
repeat_L_region	/note="M1r1 repeat: matches 3. .402 of consensus"
repeat_L_region	6989. .7284
repeat_L_region	/note="A1u5 repeat: matches 1. .296 of consensus"
repeat_L_region	8131. .8228
repeat_L_region	/note="MIR repeat: matches 124. .256 of consensus"
repeat_L_region	8437. .8740
repeat_L_region	/note="A1u5 repeat: matches 1. .304 of consensus"
repeat_L_region	8936. .9019
repeat_L_region	/note="L2 repeat: matches 2660. .2749 of consensus"
repeat_L_region	9150. .9379
repeat_L_region	/note="MIR repeat: matches 24. .259 of consensus"
repeat_L_region	10513. .10672
repeat_L_region	/note="MIR repeat: matches 63. .218 of consensus"
repeat_L_region	10674. .10795
repeat_L_region	/note="A1u5g/x repeat: matches 172. .293 of consensus"
repeat_L_region	11633. .11896
repeat_L_region	/note="L2 repeat: matches 2414. .2682 of consensus"
repeat_L_region	11870. .12134
repeat_L_region	/note="MIR repeat: matches 8. .261 of consensus"
repeat_L_region	12183. .12502

```

repeat_region /note="HALL1 repeat: matches 457. .940 of consensus"
27839. .28045
/note="LIMB8 repeat: matches 5813. .6036 of consensus"
repeat_region 28046. .28351
/note="ALUSg repeat: matches 1. .296 of consensus"
repeat_region 28352. .28494
/note="LIMB8 repeat: matches 6036. .6173 of consensus"
repeat_region 28560. .28666
/note="MIR repeat: matches 34. .142 of consensus"
repeat_region 28667. .28722
/note="FLAM.C repeat: matches 1. .125 of consensus"
repeat_region 28793. .28919
/note="MIR repeat: matches 142. .262 of consensus"
repeat_region 28887. .29074
/note="MIR repeat: matches 1. .179 of consensus"
repeat_region 29183. .29224
/note="THEIC repeat: matches 328. .371 of consensus"
repeat_region 29225. .29527
/note="ALUS repeat: matches 1. .303 of consensus"
repeat_region 29528. .29874
/note="THEIC repeat: matches 1. .328 of consensus"
repeat_region 30638. .31493
/note="L2 repeat: matches 1767. .2680 of consensus"
repeat_region 33042. .33246
/note="L2 repeat: matches 2433. .2613 of consensus"

alignment_scores:
Quality: 64.00 Length: 13
Ratio: 5.818 Gaps: 0
Percent Similarity: 84.615 Percent Identity: 53.846

alignment_block:
US-09-444-281-35 x AL162423/rev ..

Align seg 1/1 to reverse of: AL162423 from: 1 to: 88323

1 lleleulyslystrprtrpttrpttrparqrls 13
47969 TTGGTGAGCAATGGCCATGCTGGCGCATGAGCCGGAAG 47931

seq_name: gb_htg:AL356097_0

seq_documentation_block:
WPCOMMENT
Sequence split into 4 fragments LOCUS AL356097 Accession AL356097
Fragment Name Begin End
AL356097_0 1 110000
AL356097_1 100001 210000
AL356097_2 200001 310000
AL356097_3 300001 350026
LOCUS AL356097 355026 bp DNA HTG 07-APR-2001
DEFINITION Homo sapiens chromosome 1 clone RP11-180A14, *** SEQUENCING IN
PROGRESS ***, 38 unordered pieces.
AL356097 AC058797
AL356097.11 GI:11414557
HTG: HTGS_PHASE1; HTGS_DRAFT.
human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 355026)
Plumb,B.
Direct Submission
Submitted (07-APR-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk Clone
requests: clonerequests@sanger.ac.uk
On May 15, 2001 this version replaced gi:8077023
gi:9797422.
----- Genome Center
Center: Sanger Centre
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: humquerry@sanger.ac.uk

```


----- Project Information
Center project name: ba180A14
----- Summary Statistics
Assembly program: XGAP4; version 4.5
Sequencing vector: M13; M7815; 43% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Consensus quality: 33302 bases at least Q40
Consensus quality: 341682 bases at least Q30
Consensus quality: 346727 bases at least Q20
Insert size: 351326; sum-of-configs
Insert size: 176896; 2.8% error; agarose-fp
Quality coverage: 3.95x in Q20 bases; sum-of-configs Quality
coverage: 8.06x in Q20 bases; agarose-fp

Draft Sequence Produced by Whitehead Institute/MIT Center for
Genome Research, 320 Charles Street,
Cambridge, MA 02141, USA
http://www-seq.wi.mit.edu.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 38 configs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the configs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
1 16446: config of 16446 bp in length
16447 16546: gap of 100 bp
16547 22746: config of 6200 bp in length
22747 22846: gap of 100 bp
22847 45482: config of 22636 bp in length
45483 45582: gap of 100 bp
45583 49251: config of 3663 bp in length
49252 49351: gap of 100 bp
49352 53239: config of 3888 bp in length
53240 53339: gap of 100 bp
53340 58178: config of 4839 bp in length
58179 58278: gap of 100 bp
58279 71754: config of 13476 bp in length
71755 71854: gap of 100 bp
71855 76641: config of 4787 bp in length
76642 76741: gap of 100 bp
76742 80415: config of 3674 bp in length
80416 80515: gap of 100 bp
80516 89902: config of 9387 bp in length
89903 90002: gap of 100 bp
90003 93661: config of 3659 bp in length
93662 93761: gap of 100 bp
93762 101785: config of 8024 bp in length
101786 101885: gap of 100 bp
101886 109333: config of 7448 bp in length
109334 109433: gap of 100 bp
109434 121288: config of 11855 bp in length
121289 121388: gap of 100 bp
121389 128767: config of 7379 bp in length
128768 128867: gap of 100 bp
128868 131917: config of 3050 bp in length
131918 132017: gap of 100 bp
132018 138795: config of 6778 bp in length
138796 138895: gap of 100 bp
138896 144660: config of 5765 bp in length
144661 144760: gap of 100 bp
144761 156587: config of 11827 bp in length
156588 156687: gap of 100 bp
156688 158939: config of 2252 bp in length
158940 161353: gap of 100 bp
161354 161453: config of 2314 bp in length
161454 165170: config of 3717 bp in length
165171 165270: gap of 100 bp
165271 168479: config of 3209 bp in length
168480 168579: gap of 100 bp

168580 172188: config of 3609 bp in length
172189 172288: gap of 100 bp
172289 177084: config of 4796 bp in length
177085 177184: gap of 100 bp
177185 199876: config of 22692 bp in length
199877 199976: gap of 100 bp
199977 222186: config of 22210 bp in length
222187 222286: gap of 100 bp
222287 240235: config of 17949 bp in length
240236 240335: gap of 100 bp
240336 263715: config of 23360 bp in length
263716 263815: gap of 100 bp
263816 269270: config of 5455 bp in length
269271 269370: gap of 100 bp
269371 271745: config of 2375 bp in length
271746 271845: gap of 100 bp
271846 277024: config of 5179 bp in length
277025 277124: gap of 100 bp
277125 280037: config of 2913 bp in length
280038 280137: gap of 100 bp
280138 307328: config of 27191 bp in length
307329 307428: gap of 100 bp
307429 313985: config of 6557 bp in length
313986 314085: gap of 100 bp
314086 330085: config of 16000 bp in length
330086 330185: gap of 100 bp
330186 334887: config of 4702 bp in length
334888 334987: gap of 100 bp
334988 355026: config of 20039 bp in length.
Location/Qualifiers
1. .355026
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="1"
/clone="RP11-180A14"
/clone_lib="RPC1-11.1"
1. .16446
/note="assembly_fragment:02644
fragment_chain:1"
16547. .22746
/note="assembly_fragment:02589
fragment_chain:1"
22847. .45482
/note="assembly_fragment:03035
fragment_chain:1"
45583. .49251
/note="assembly_fragment:00032"
49352. .53239
/note="assembly_fragment:00060"
53340. .58178
/note="assembly_fragment:00105"
58279. .71754
/note="assembly_fragment:00250"
71855. .76641
/note="assembly_fragment:00381"
76742. .80415
/note="assembly_fragment:00389"
80516. .89902
/note="assembly_fragment:00591"
90003. .93661
/note="assembly_fragment:00750"
93762. .101785
/note="assembly_fragment:00754"
101886. .109333
/note="assembly_fragment:00819"
109434. .121288
/note="assembly_fragment:00889"
121389. .128767
/note="assembly_fragment:00932"
128868. .131917
/note="assembly_fragment:00972"
132018. .138795
/note="assembly_fragment:01033"


```

* 91004 91103: gap of 100 bp
* 91104 109428: contig of 18325 bp in length
* 109429 109528: gap of 100 bp
* 109529 135277: contig of 25749 bp in length
* 135278 135377: gap of 100 bp
* 135378 159391: contig of 24014 bp in length.
Location/Qualifiers
1. 159391

```

FEATURES

source

```

/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="5"
/map="5"
/clone="RP11-67212"
/clone.lib="RPC1-11 Human Male BAC"
1. 1373
misc_feature
/note="assembly_fragment"
1474. 3371
misc_feature
/note="assembly_fragment"
3472. 5991
misc_feature
/note="assembly_fragment"
6092. 8001
misc_feature
/note="assembly_fragment"
8102. 10452
misc_feature
/note="assembly_fragment"
10553. 13499
misc_feature
/note="assembly_fragment"
13600. 17300
misc_feature
/note="assembly_fragment"
17401. 23167
misc_feature
/note="assembly_fragment"
23268. 27394
misc_feature
/note="assembly_fragment"
clone_end:7
vector_side:right"
27495. 31941
misc_feature
/note="assembly_fragment"
32042. 36867
misc_feature
/note="assembly_fragment"
36968. 43237
misc_feature
/note="assembly_fragment"
43338. 52793
misc_feature
/note="assembly_fragment"
52896. 61446
misc_feature
/note="assembly_fragment"
61547. 68822
misc_feature
/note="assembly_fragment"
68923. 79907
misc_feature
/note="assembly_fragment"
80008. 91003
misc_feature
/note="assembly_fragment"
91104. 109428
misc_feature
/note="assembly_fragment"
109529. 135277
misc_feature
/note="assembly_fragment"
135378. 159391
misc_feature
/note="assembly_fragment"
clone_end:SP6
vector_side:right"
BASE COUNT 43727 a 35737 c 36282 g 41744 t 1901 others
ORIGIN

```

```

alignment_scores:
Quality: 64.00 Length: 9
Ratio: 7.11 Gaps: 0
Percent similarity: 100.000 Percent identity: 77.778

```

alignment_block:

US-09-444-281-35 x AC027113

Align seg 1/1 to: AC027113 from: 1 to: 159391

5 TrpProTPrTrpPrpArGArGlys 13

87682 TGGCCCTGGTGGCTTGGAGGAGGAG 87708

seq_name: gb_hgt:AL358473

seq_documentation_block:

```

LOCUS AL358473 187845 bp DNA HTG 07-APR-2001
DEFINITION Homo sapiens chromosome 1 clone RP11-168016, *** SEQUENCING IN
PROGRESS *** 7 unordered pieces.
ACCESSION AL358473 AC027712
VERSION AL358473.8 GI:11322009
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 187845)
McLay, K.
Direct Submission
Submitted (06-APR-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk
requests: clonerequest@sanger.ac.uk
On May 15, 2001 this sequence version replaced gi:7712176
gi:10800605.
----- Genome Center
Center: Sanger Centre
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: humquerry@sanger.ac.uk
----- Project Information
Center project name: BA168016
----- Summary Statistics
Assembly program: XGAP4; version 4.5
Sequencing vector: M13; M77815; 36% of reads
Sequencing vector: plasmid; L08752; 63% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Consensus quality: 186132 bases at least Q40
Consensus quality: 186498 bases at least Q30
Consensus quality: 186851 bases at least Q20
Insert size: 187245; sum-of-contigs
Insert size: 186146; agarose-fp
Quality coverage: 10.30x in Q20 bases; sum-of-contigs Quality
coverage: 10.58x in Q20 bases; agarose-fp
-----
Draft Sequence Produced by Whitehead Institute/MIT Center for
Genome Research, 320 Charles Street,
Cambridge, MA 02141, USA
http://www-seq.wi.mit.edu.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 7 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
1 11961: contig of 11961 bp in length
* 11962 12061: gap of 100 bp
* 12062 95058: contig of 82997 bp in length
* 95059 95158: gap of 100 bp
* 95159 124623: contig of 29465 bp in length
* 124624 124723: gap of 100 bp
* 124724 128609: contig of 4886 bp in length
* 128610 129709: gap of 100 bp
* 129710 151381: contig of 21672 bp in length
* 151382 151481: gap of 100 bp
* 151482 174927: contig of 23446 bp in length
* 174928 175027: gap of 100 bp
* 175028 187845: contig of 12818 bp in length.
Location/Qualifiers
1. 187845
/organism="Homo sapiens"
/db_xref="taxon:9606"

```

FEATURES

source

```

misc_feature /chromosome="1"
/clone="RP11-168016"
/clone_lib="RPC1-11.1"
1..11961
/note="assembly_fragment:01682
fragment_chain:1
clone_end:SP6
vector_side:left"
misc_feature 12062..95058
/note="assembly_fragment:01450
fragment_chain:1"
misc_feature 95159..124623
/note="assembly_fragment:01175
fragment_chain:1"
misc_feature 124724..129609
/note="assembly_fragment:04318
fragment_chain:1"
misc_feature 129710..151381
/note="assembly_fragment:00767
fragment_chain:2"
misc_feature 151482..174927
/note="assembly_fragment:01178
fragment_chain:2"
misc_feature 175028..187845
/note="assembly_fragment:02771
clone_end:R7
vector_side:right"
BASE COUNT 46168 a 48356 c 47958 g 44763 t 600 others
ORIGIN

```

```

alignment_scores:      Quality: 64.00      Length: 9
Ratio: 7.111          Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

```

alignment_block:
US-09-444-281-35 x AL358473

Align seg 1/1 to: AL358473 from: 1 to: 187845

```

5 TrpProTrrPrpTrrPrpAtrGlys 13
|||||
133213 TGGCCCTGTGGCTTGGAGGAGGAG 133239

```


25 ATCTGAAAAATGCGCTGGTGGCGCTGCGCTGATAA 63
seq_name: /SIDS8/gcgdata/geneseq/geneseqn/NA2000.DAT:AAA27291
seq_documentation_block:
ID AAA27291 standard; DNA: 114 BP.
XX
AC AAA27291;
XX
DT 20-SEP-2000 (first entry)
XX
DE Oligonucleotide used for synthesis of MBI-11 fragment.
XX
KW Oligonucleotide; cellulose binding domain; CBD; cationic peptide;
XX MBI-11; indolicidin; bovine; ss.
XX
OS Synthetic.
XX
PN WO200031279-A2.
XX
PD 02-JUN-2000.
XX
PF 19-NOV-1999; 99WO-CA01107.
XX
PR 20-NOV-1998; 98US-0109218.
XX
PA (MICR-) MICROLOGIX BIOTECH INC.
XX
PI Burian J, Bartfeld D;
XX
DR WPI: 2000-400086/34.
XX
PT Multi-domain fusion protein expression cassette used for high yield
XX stable production of foreign peptide gene products -
XX
PS Example 4; Page 37; 73pp; English.
XX
CC A novel method allows the efficient production of cationic peptides in
CC recombinant host cells. The method involves construction of a
CC multi-domain fusion protein expression cassette comprising a promoter and
CC a nucleic acid molecule expressed as an insoluble protein. The inclusion
CC of anionic peptide sequences in the linker sequences neutralises the
CC positive charge of the cationic peptide so that the charge of the
CC fusion protein is controlled. This cassette allows high yield, stable
CC production of the cationic peptide. Cationic peptides such as
CC bovine indolicidin may be used as antimicrobial agents. The present
CC sequence is an oligonucleotide that was used to synthesise a
CC MBI-11 fragment. MBI-11 is a cationic peptide derived from modifications
CC of indolicidin.
XX
SQ Sequence 114 BP; 25 A; 26 C; 30 G; 33 T; 0 other;

alignment_scores:
Quality: 91.00 Length: 13
Ratio: 7.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x AAA27291
Align seg 1/1 to: AAA27291 from: 1 to: 114

1 11leuLysLysTrpProTrpTrpProTrpArgArgLys 13
|||||
41 ATCTGAAAAATGCGCTGGTGGCGCTGCGCTGATAA 79

seq_name: /SIDS8/gcgdata/geneseq/geneseqn/NA2000.DAT:AAA27296
seq_documentation_block:
ID AAA27296 standard; DNA: 108 BP.
XX
AC AAA27296;
XX

XX
DT 20-SEP-2000 (first entry)
XX
DE Oligonucleotide used for synthesis of MBI 2X11B7 poly cassette.
XX
KW Oligonucleotide; cellulose binding domain; CBD; cationic peptide;
XX MBI-11; indolicidin; bovine; ss.
XX
OS Synthetic.
XX
PN WO200031279-A2.
XX
PD 02-JUN-2000.
XX
PF 19-NOV-1999; 99WO-CA01107.
XX
PR 20-NOV-1998; 98US-0109218.
XX
PA (MICR-) MICROLOGIX BIOTECH INC.
XX
PI Burian J, Bartfeld D;
XX
DR WPI: 2000-400086/34.
XX
PT Multi-domain fusion protein expression cassette used for high yield
XX stable production of foreign peptide gene products -
XX
PS Example 5; Page 39; 73pp; English.
XX
CC A novel method allows the efficient production of cationic peptides in
CC recombinant host cells. The method involves construction of a
CC multi-domain fusion protein expression cassette comprising a promoter and
CC a nucleic acid molecule expressed as an insoluble protein. The inclusion
CC of anionic peptide sequences in the linker sequences neutralises the
CC positive charge of the cationic peptide so that the charge of the
CC fusion protein is controlled. This cassette allows high yield, stable
CC production of the cationic peptide. Cationic peptides such as
CC bovine indolicidin may be used as antimicrobial agents. The present
CC sequence is an oligonucleotide that was used to synthesise a
CC MBI-11B7 fragment. This fragment was used in the expression
CC cassette. MBI-11B7 is a cationic peptide derived from modifications
CC of indolicidin.
XX
SQ Sequence 108 BP; 18 A; 33 C; 31 G; 26 T; 0 other;

alignment_scores:
Quality: 76.00 Length: 13
Ratio: 6.333 Gaps: 0
Percent Similarity: 92.308 Percent Identity: 69.231

alignment_block:
US-09-444-281-35 x AAA27296
Align seg 1/1 to: AAA27296 from: 1 to: 108

1 11leuLysLysTrpProTrpTrpProTrpArgArgLys 13
|||||
38 ATGATTCGTCGTTGGCGCTGGCGCTGCGCTGCGCAA 76

seq_name: /SIDS8/gcgdata/geneseq/geneseqn/NA2000.DAT:AAA27298
seq_documentation_block:
ID AAA27298 standard; DNA: 114 BP.
XX
AC AAA27298;
XX
DT 20-SEP-2000 (first entry)
XX
DE Oligonucleotide used for synthesis of MBI 11B7 first cassette.
XX
KW Oligonucleotide; cellulose binding domain; CBD; cationic peptide;
XX MBI-11; indolicidin; bovine; ss.

```

OS Synthetic.
XX
PN W0200031279-A2.
XX
PD 02-JUN-2000.
XX
PF 19-NOV-1999; 99W0-CA01107.
XX PR 20-NOV-1998; 98US-0109218.
XX PA (MICR-) MICROLOGIX BIOTECH INC.
XX PI Burian J, Bartfeld D;
XX WP1: 2000-400086/34.
XX
PT Multi-domain fusion protein expression cassette used for high yield
PR stable production of foreign peptide gene products -
XX
PS Example 5; Page 40; 73pp; English.
XX
CC A novel method allows the efficient production of cationic peptides in
CC recombinant host cells. The method involves construction of a
CC multi-domain fusion protein expression cassette comprising a promoter and
CC a nucleic acid molecule expressed as an insoluble protein. The inclusion
CC of anionic peptide sequences in the linker sequences neutralises the
CC positive charge of the cationic peptide so that the charge of the
CC fusion protein is controlled. This cassette allows high yield, stable
CC production of the cationic peptide. Cationic peptides such as
CC bovine indolicidin may be used as antimicrobial agents. The present
CC sequence is an oligonucleotide that was used to synthesise a
CC MBI-11B7 fragment. This fragment was used in the expression
CC cassette. MBI-11B7 is a cationic peptide derived from modifications
CC of indolicidin.
XX
SQ Sequence 114 BP; 20 A; 34 C; 32 G; 28 T; 0 other:
XX
alignment_scores:
Quality: 76.00 Length: 13
Ratio: 6.333 Gaps: 0
Percent Similarity: 92.308 Percent Identity: 69.231
XX
alignment_block:
US-09-444-281-35 x AAA27298*
XX
Align seg 1/1 to: AAA27298 from: 1 to: 114
XX
1 lileulyslstprptprtprtpatargalys 13
XXXX : : : : :
44 ATGATTCGTGGCGTGTGGCGTGCGTCGCAGAA 82
XX
seq_name: /SIDSB/gcgdata/geneseq/geneseqn/NA2000.DAT:AAA27294
XX
seq_documentation_block:
ID AAA27294 standard; DNA; 451 BP.
XX
AC AAA27294:
XX
DT 20-SEP-2000 (first entry)
XX
DE Oligonucleotide used for synthesis of MBI 2X11B7 last cassette.
XX
KW Oligonucleotide; cellulose binding domain; CBD; cationic peptide;
XX MBI-11; indolicidin; bovine; ss.
XX
OS Synthetic.
XX
PN W0200031279-A2.
XX
PD 02-JUN-2000.

```

```

PE 19-NOV-1999; 99WO-CA01107.
XX
PR 20-NOV-1998; 98US-0109218.
XX
PA (MICR-) MICROLOGIX BIOTECH INC.
XX
PI Burian J, Bartfeld D:
XX
DR WPI; 2000-40086/34.
XX
PT Multi-domain fusion protein expression cassette used for high yield
PT stable production of foreign peptide gene products -
XX
PS Example 5; Page 38; 73pp: English.
XX
CC A novel method allows the efficient production of cationic peptides in
CC recombinant host cells. The method involves construction of a
CC multi-domain fusion protein expression cassette comprising a promoter and
CC a nucleic acid molecule expressed as an insoluble protein. The inclusion
CC of anionic peptide sequences in the linker sequences neutralises the
CC positive charge of the cationic peptide so that the charge of the
CC fusion protein is controlled. This cassette allows high yield, stable
CC production of the cationic peptide. Cationic peptides such as
CC bovine indolicidin may be used as antimicrobial agents. The present
CC sequence is an oligonucleotide that was used to synthesise a
CC MB1-11B7 fragment. This fragment was used in the expression
CC cassette. MB1-11B7 is a cationic peptide derived from modifications
CC of indolicidin.
XX
SQ Sequence 151 BP; 22 A; 44 C; 49 G; 36 T; 0 other;

Alignment_scores:
    Quality: 76.00      Length: 13
    Ratio: 6.333      Gaps: 0
    Percent Similarity: 92.308      Percent Identity: 69.231

Alignment_block:
US-09-444-281-35 x AAA27294 ..

Align seg 1/1 to: AAA27294 from: 1 to: 151

1 ILELeuLysLysTTPProTTPTrPProTTPArgArgLys 13
   :::::  ::::::::::::::::::::::::::::
38 ATGATTCTCGCGTTGGCCCTGTGGTGGCCGTGGGTCGCAAA 76

seq_name: /SID58/gcgdata/geneseq/geneseqn/AA1999.DAT:AAV83788

seq_documentation_block:
ID AAV83788 standard; DNA; 39 BP.
XX
AC AAV83788;
XX
DT 19-MAR-1999 (first entry)
XX
DE Antimicrobial peptide Indolicidin encoding DNA.
XX
KW Antimicrobial; fusion; acidic peptide; recombinant; microorganism;
KW guaneterin; basic peptide; Indolicidin; ss.
XX
OS Synthetic.
OS Bos sp.
XX
FH Key
FH CDS
FT 1.39
FT /tag= a
FT /note= "The start and stop codons are not indicated"
XX
PN WO9854336-A1.
XX
DD 03-DEC-1998.
XX
PP 28-MAY-1998; 98WO-KR00132.

```

XX 09-APR-1998; 98KR-0013372.
PR 28-MAY-1997; 97KR-0021312.
XX
XX (KOAD) KOREA ADV INST SCI & TECHNOLOGY.
PA (SAMY-) SAMYANG GENEX CORP.
XX
PI Hong S, Kang MH, Kim JH, Kim S, Lee H, Lee JH;
XX WPI, 1999-059844/05.
DR P-PSDB; AAW87609.
XX
PT New method for mass production of antimicrobial peptides - by
PT constructing fusion genes comprising acidic and antimicrobial
PT peptide genes and transforming host with vector containing these
PS
XX Example 6; Page 18; 52pp; English.
XX
CC The invention relates to mass production of antimicrobial peptides. The
CC method comprises constructing a fusion gene containing a first gene
CC encoding a negatively charged acidic peptide having at least two cysteine
CC residues, and a second gene encoding a positively charged basic
CC antimicrobial peptide. A host microorganism is transformed with a vector
CC containing the fusion gene and then cultured. The expressed antimicrobial
CC peptide is then recovered. The method is used to mass produce
CC antimicrobial peptides in recombinant microorganisms. The inhibitory
CC effect of the expressed antimicrobial peptide upon the growth of the host
CC microorganism is considerably reduced by fusing it to the acidic peptide.
CC Therefore, the use of the fusion gene provides an economic, recombinant
CC alternative of mass producing antimicrobial peptides, which overcomes the
CC disadvantages of low productivity and poor economy, previously
CC encountered by recombinant and chemical methods. The present sequence
CC represents the DNA encoding an antimicrobial peptide indolicidin. This
CC can be used along with the acidic peptide guanemycin gene in the
CC construction of the fusion gene.
XX
SQ Sequence 39 BP; 4 A; 10 C; 16 G; 9 T; 0 other;

alignment_scores:
Quality: 73.00 Length: 9
Ratio: 8.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x AAV83788 ..
Align seg 1/1 to: AAV83788 from: 1 to: 39

4 LysTrpProTfRTPProTfRPaRgArg 12
|||||
13 AATGCGCGTGTGCGCGTGTGCGTGTGCT 39

seq_name: /SID8/gcgdata/geneseq/geneseqn/NA2000.DAT:AAZ29389

seq_documentation_block:
ID AAZ29389 standard; DNA; 47 BP.
XX
AC AAZ29389;
XX
DT 29-FEB-2000 (first entry)
XX
DE PCR primer-15 for synthesis of antimicrobial peptide indolicidin gene.
XX
KW PCR primer; anti-microbial peptide; indolicidin gene; DNA construct;
KW glutamine pyrophosphoribosyl pyrophosphate amidotransferase gene;
KW purf gene; fusion peptide; mass production; pharmaceutical industry;
KW food industry; ss.
XX
OS Synthetic.
XX
XX MO9964611-A1.
PN
XX

PD 16-DEC-1999.
XX
XX 08-JUN-1999; 99MO-KR00282.
PF
XX
XX 09-JUN-1998; 98KR-0022117.
PR 14-MAY-1999; 99KR-0017920.
XX
XX (SAMY-) SAMYANG GENEX CORP.
PA
PI Kim JH, Kang MH, Lee J, Park SH, Lee JW, Hong SS, Lee H;
XX WPI, 2000-097542/08.
DR
XX
XX
PT New DNA constructs useful for mass production of antimicrobial peptides
PT in microorganism hosts -
XX
XX
PS Example 1; Page 13; 67pp; English.
XX
XX
CC The present sequence is a chemically synthesised PCR primer which was
CC used to synthesise a gene encoding antimicrobial peptide indolicidin.
CC The antimicrobial peptide gene is used in a DNA construct that comprises
CC entire, partial or a derivative of purf gene (glutamine
CC pyrophosphoribosyl pyrophosphate amidotransferase gene). The DNA
CC construct allows mass production of the antimicrobial peptide in
CC microbial hosts without killing the host cells. The antimicrobial
CC peptides are useful commercially in the pharmaceutical and
CC food industries.
XX
SQ Sequence 47 BP; 6 A; 11 C; 19 G; 11 T; 0 other;

alignment_scores:
Quality: 73.00 Length: 9
Ratio: 8.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x AAZ29389 ..
Align seg 1/1 to: AAZ29389 from: 1 to: 47

4 LysTrpProTfRTPProTfRPaRgArg 12
|||||
17 AATGCGCGTGTGCGCGTGTGCGTGTGCT 43

seq_name: /SID8/gcgdata/geneseq/geneseqn/NA2000.DAT:AAZ29390

seq_documentation_block:
ID AAZ29390 standard; DNA; 47 BP.
XX
AC AAZ29390;
XX
DT 29-FEB-2000 (first entry)
XX
DE PCR primer-16 for synthesis of antimicrobial peptide indolicidin gene.
XX
XX
KW PCR primer; anti-microbial peptide; indolicidin gene; DNA construct;
KW glutamine pyrophosphoribosyl pyrophosphate amidotransferase gene;
KW purf gene; fusion peptide; mass production; pharmaceutical industry;
KW food industry; ss.
XX
OS Synthetic.
XX
XX MO9964611-A1.
PN
XX
XX 16-DEC-1999.
PD
XX
PF 08-JUN-1999; 99MO-KR00282.
XX
XX
PR 09-JUN-1998; 98KR-0022117.
PR 14-MAY-1999; 99KR-0017920.
XX
XX (SAMY-) SAMYANG GENEX CORP.
PA


```

PI Kim JH, Kang MH, Lee J, Park SH, Lee JW, Hong SS, Lee H;
XX
XX WPI: 2000-097542/08.
XX
XX New DNA constructs useful for mass production of antimicrobial peptides
XX in microorganism hosts -
XX
XX Example 1; Page 13; 67pp: English.
XX
XX The present sequence is a chemically synthesised PCR primer which was
CC used to synthesise a gene encoding antimicrobial peptide Indolicidin.
CC The antimicrobial peptide gene is used in a DNA construct that comprises
CC entire, partial or a derivative of purF gene (glutamine
CC pyrophosphoribosyl pyrophosphate amidotransferase gene). The DNA
CC construct allows mass production of the antimicrobial peptide in
CC microbial hosts without killing the host cells. The antimicrobial
CC peptides are useful commercially in the pharmaceutical and
CC food industries.
XX
XX Sequence 47 BP; 12 A; 18 C; 10 G; 7 T; 0 other;
SQ
alignment_scores:
Quality: 73.00 Length: 9
Ratio: 8.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000
alignment_block:
US-09-444-281-35 x AAZ29390/rev
Align seg 1/1 to reverse of: AAZ29390 from: 1 to: 47
4 LysrrpProtrTPrrProtrPArgarg 12
|||||
35 AAATGGCCGTGGTGGCCGTGGCGTGTGT 9
seq_name: /SID58/gcgdata/geneseq/geneseqn/NA2000.DAT:AAZ29364
seq_documentation_block:
ID AAZ29364 standard; DNA: 53 BP.
AC AAZ29364:
XX
XX 29-FEB-2000 (first entry)
XX
XX Antimicrobial peptide, Indolicidin encoding DNA.
DE
XX
XX purF gene; glutamine pyrophosphoribosyl pyrophosphate amidotransferase;
KW purF derivative; fusion partner; antimicrobial peptide; Indolicidin;
KW mass production; cleavage site; hydroxylamine; CNBR; DNA construct; cow;
XX neutralise; toxicity; pharmaceutical industry; food industry; ds.
XX
XX Bos taurus.
OS
XX
XX Key Location/Qualifiers
XX FH 5..46
XX CDS FT /*tag= "B"
XX FT /product= "Indolicidin peptide"
XX FT /note= "Antimicrobial peptide used in DNA construct"
XX
XX WO964611-A1.
XX
XX 16-DEC-1999.
XX
XX 08-JUN-1999; 99WO-KR00282.
XX
XX 09-JUN-1998; 98KR-0022117.
XX 14-MAY-1999; 99KR-0017920.
XX
XX (SAMV-) SAMYANG GENEX CORP.
XX
XX Kim JH, Kang MH, Lee J, Park SH, Lee JW, Hong SS, Lee H;
XX

```

```

XX  WPI: 2000-097542/08.
DR  P-PSDB: AAY44324.
XX
PT  New DNA constructs useful for mass production of antimicrobial peptides
PT  in microorganism hosts -
XX
PS  Claim 1; Fig 1; 67pp; English.
XX
CC  The present DNA sequence encodes an antimicrobial peptide, indolicidin
CC  derived from cow, Bos taurus. It is used along with a derivative
CC  of purf gene sequence that functions as a fusion partner.
CC  A DNA construct that comprises, this antimicrobial peptide encoding
CC  sequence and the entire, partial or derivative of purf gene, is used for
CC  mass production of the antimicrobial peptide in microorganisms without
CC  killing the host cells. Use of the purf gene derivative sequence,
CC  neutralises the toxicity of the antimicrobial peptides against the host
CC  microorganism. The antimicrobial peptides are useful commercially in the
CC  pharmaceutical and food industries.
XX
SO  Sequence 53 BP; 8 A; 12 C; 20 G; 13 T; 0 other;

alignment_scores:
    quality: 73.00      Length: 9
    Ratio: 8.111      Gaps: 0
    Percent Similarity: 100.000      Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x AA229364 ..

Align seg 1/1 to: AA229364 from: 1 to: 53

      4 LYSTRPpROTPRTpPpOTpPARGArg 12
      ||| ||||| ||||| ||||| |||||
      17 AAATGCGCCGTGGTGGCCGTGGCGTGTGT 43      *

seq_name: /SIDSB/gcgdata/geneseq/geneseqn/NA2000.DAT:AA240246
seq_documentation_block:
ID  AA240246 standard; DNA: 69 BP.
XX
AC  AA240246;
XX
DT  23-FEB-2000 (first entry)
XX
DE  Oligonucleotide for cloning indolicidin peptide coding sequence.
XX
KW  Indolicidin; bactericin; sulphate-reducing bacteria; growth inhibitor;
KW  corrosion; degradation; metal; concrete; cement; dental implant; biofilm
XX  ss.
XX
OS  Synthetic.
OS  Bacillus sp.
XX
PN  W09956553-A1.
XX
PD  11-NOV-1999.
XX
PF  03-MAY-1999; 99WO-US09675.
XX
PR  06-MAY-1998; 98US-0074037.
PR  31-MAR-1999; 99US-0282277.
XX
PA  (REGC ) UNIV CALIFORNIA.
XX
PI  Wood TK, Jayaraman A, Earhman JC;
XX
DR  WPI: 2000-052882/04.
XX
PT  Inhibiting growth of sulphate-reducing bacteria using other bacteria,
PT  particularly for protection of metals and concrete -
XX

```

PS Example 4; Fig 1; 84pp; English.

XX This sequence represents an oligonucleotide for cloning the non-amidated
CC indolicidin peptide coding sequence. The invention relates to a method
CC for inhibiting growth of sulphate-reducing bacteria (A) on a material (B)
CC sensitive to corrosion or degradation, by applying to (B) a bacterium (C)
CC that secretes a compound (I) able to inhibit growth of (A). The method is
CC used to protect metal, concrete or cement against corrosion and
CC degradation, but (B) can also be used to protect dental implants. (B) is
CC present in an open or closed system (e.g. water cooling tower, liquid
CC storage container, fuel tank, sewer or drainage system etc.) or part of a
CC bridge or other structure. The method is more effective and less
CC expensive than known methods for inhibiting (A), and reduces the amount
CC of toxic chemicals released. Conventional biofilms of aerobic organisms
CC tend to encourage growth of (A), and addition of (C) to the biofilm
CC prevents this. A single application of (C) lasts for a long time, and (I)
CC are produced exactly where they are required and inhibit (A) without
CC significant impact on other organisms (this effect includes reducing
CC resistance of (A) to conventional biocides, which may then be used in
CC reduced amounts). If local damage to the biofilm occurs, the underlying
CC material is still protected by diffusion of (I) from neighbouring areas.

XX Sequence 69 BP; 14 A; 18 C; 20 G; 17 T; 0 other;

XX SQ

alignment_scores:
Quality: 73.00 Length: 9
Ratio: 8.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x AA240246

Align seg 1/1 to: AA240246 from: 1 to: 69

4 LysTrpProTrrPrProTrrPargArg 12
|||||
28 AATGGCGCTGGTGGCGCTGGCGCGC 54

seq_name: /SID58/gcdata/geneseq/geneseqn/NA2000.DAT:AA249764

seq_documentation_block:
ID AA249764 standard: DNA; 211 BP.

XX AC AA249764;
XX DF 18-APR-2000 (first entry)
XX DE Poly-(Indol (1-13)-Met-Ala-Arg-Ile-Ala-Met)₃ DNA.
XX KW Crosslinked indolicidin analog: X-indolicidin; poly-indol 1-13;
KW stability: bovine neutrophil; antimicrobial; antibacterial; fungicide;
KW protozoacide; virucide; anti-HIV; human immunodeficiency virus-1;
KW HIV-1; gram positive bacteria; gram negative; Staphylococcus aureus;
KW Escherichia coli; Salmonella typhimurium; yeast; fungi; protozoa;
KW Candida albicans; Cryptococcus neoformans; Giardia; Acanthamoeba;
KW hexapeptide spacer; ds.
XX OS Synthetic.
OS Bos sp.
XX Key CDS
FH Location/Qualifiers
FT 8..199
FT /*tag= a
FT /product= "Poly-(indol(1-13)-Met-Ala-Arg-Ile-Ala-Met)₃"
FT /note= "encodes three copies of indol 1-13, each
FT separated by Met-Ala-Arg-Ile-Ala-Met spacer sequence"
FT 1..21
FT /*tag= b
FT primer_bind complement (191..211)
FT /*tag= c
FT 68..71
FT misc_feature
FT /*tag= d

FT /*note= "corresponds to overlap in oligonucleotides
FT used for ligation"
FT 148..151
FT /*tag= e
FT /note= "corresponds to overlap in oligonucleotides
FT used for ligation"

XX W09965510-A1.
XX PD 23-DEC-1999.
XX PF 20-MAY-1999; 99W0-US11165.
XX PR 18-JUN-1998; 98US-0099631.
XX PA (REGC) UNIV CALIFORNIA.
XX PI Selsted ME, Osapay K;
XX WP1: 2000-147133/13.
XX DR P-PSDB: AAY44668.
XX XX

XX The patent discloses crosslinked analogs of indolicidin (indol 1-13)
XX which is a naturally occurring peptide isolated from bovine neutrophils
XX and has antimicrobial activity. The crosslinked indolicidin
XX (X-indolicidin) analogs are stable and have antimicrobial activity
XX against gram positive and negative bacteria (e.g. Staphylococcus aureus,
XX Escherichia coli and Salmonella typhimurium), yeasts and fungi (e.g.
XX Candida albicans, Cryptococcus neoformans), protozoa (e.g. Giardia
XX species and Acanthamoeba species), and viruses (e.g. HIV-1).
XX They can be used for reducing or inhibiting the growth or survival of
XX microorganisms in an environment e.g. a food or food product, a
XX solution, an inanimate object comprising a surface, or a mammal.
XX The present sequence is a DNA encoding a protein comprising three
XX copies of indol 1-13 each separated by a hexapeptide spacer sequence.
XX The sequence was used to produce a recombinant construct for the
XX expression of indol-homoserine (Hse) analog. The ability of
XX indol-Hse analog to maintain antimicrobial activity provides a means to
XX produce X-indolicidin analog precursors in sufficient quantities.

XX SQ Sequence 211 BP; 36 A; 50 C; 74 G; 51 T; 0 other;

alignment_scores:
Quality: 73.00 Length: 9
Ratio: 8.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x AA249764

Align seg 1/1 to: AA249764 from: 1 to: 211

4 LysTrpProTrrPrProTrrPargArg 12
|||||
38 AATGGCGCTGGTGGCGCTGGCGCTGG 64

seq_name: /SID58/gcdata/geneseq/geneseqn/NA2000.DAT:AA245123

seq_documentation_block:
ID AA245123 standard: DNA; 211 BP.

XX AC AA245123;
XX DF 28-FEB-2000 (first entry)
XX DE Indolicidin fusion peptide nucleotide sequence.
XX XX

KM Indolicidin analogue; antimicrobial activity; helminth; bacteria; virus;
KM treatment; inhibit growth; micro-organism; contact lens solution;
KM transgenic plant; surgical instrument; yeast; fungi; protozoa; ss.
XX
OS Synthetic.
XX
XX WO958141-A1.
XX
XX 18-NOV-1999.
XX
XX 05-MAY-1999; 99WO-US09942.
XX
XX 12-MAY-1998; 98US-0076227.
XX
XX (REGC) UNIV CALIFORNIA.
XX
XX Selssted ME;
XX
XX WPI: 2000-053028/04.
DR P-PSDB; AAY57142.
XX
XX New indolicidin analogues, active against bacteria, yeast, fungi,
PT protozoa and virus, used for, e.g., treating infections -
XX
XX Disclosure; Fig 6; 62pp; English.
PS
XX
XX This is the nucleotide sequence of an example of a fusion protein which
CC consists of an indolicidin analogue linked to another peptide.
CC Peptides AAY57109-157138 and AAY57143-157144 are new indolicidin
CC analogues which have a homoserine residue and/or a truncated amino
CC terminal region. The analogues have the following amino acid sequence:
CC Xaa1-Xaa2-Xaa3-Xaa4-Xaa5-Xaa6-Pro-Xaa6-Pro-Xaa6-Xaa7-Xaa7-Xaa8
CC where:
CC Xaa1 = Ile, Leu, Val, Ala, Gly or absent;
CC Xaa2 = Ile, Leu, Val, Ala, Gly or absent;
CC Xaa3 = Pro or absent;
CC Xaa4 = Trp, Phe or absent;
CC Xaa5 = Arg, Lys or absent;
CC Xaa6 = Trp or Phe;
CC Xaa7 = Arg, Lys or absent;
CC Xaa8 = homoserine (Hse), Met, Xaa9-Met or absent, and
CC Xaa9 = at least one amino acid;
CC provided that if Xaa1 is present, Xaa8 = Hse, Met or Met-Xaa9-Met;
CC and further provided that: if Xaa2 is absent, Xaa1 is absent; if Xaa3 is
CC absent, Xaa1 and Xaa2 are absent; if Xaa4 is absent, Xaa1, Xaa2 and Xaa3
CC are absent; and if Xaa5 is absent, Xaa1, Xaa2, Xaa3 and Xaa4 are absent.
CC The indolicidin analogues can be used to create a fusion polypeptide
CC consisting of the analogue linked to a peptide. The indolicidin
CC analogues have antimicrobial activity against gram positive bacteria,
CC gram negative bacteria, yeast, fungus, protozoa and viruses (e.g. HIV-1).
CC They are also active against helminths. The analogues can be used for
CC reducing or inhibiting growth or survival of a microorganism. They can be
CC used for treating infections. They can also be included in a liquid such
CC as water or an aqueous solution, e.g. contact lens solution. The
CC analogues have potential uses in food products, and in objects such as
CC the surface of an instrument used to prepare food or to perform surgery.
CC Transgenic plants or animals useful in the food industry can be produced
CC by introducing a nucleic acid molecule encoding an indolicidin analogue
CC into the germ-line cells of such organisms.
XX
XX Sequence 211 BP; 36 A; 50 C; 74 G; 51 T; 0 other;

alignment_scores:
Quality: 73.00 Length: 9
Ratio: 8.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-35 x AAZ45123

Align seg 1/1 to: AAZ45123 from: 1 to: 211

4 LysTrpProTrpTrpProTrpArgArg 12
|||||
38 AATGCGCGGTGGTGGCGCGGTGCT 64

seq_name: /SIDS8/gcgdata/geneseq/geneseqn/NA1999.DAT:AAZ20646

seq_documentation_block:

ID AAZ20646 standard; RNA: 6446 BP.

XX AAZ20646;

XX 26-NOV-1999 (first entry)

DE TMV-based virus TMV861 coat protein read-through RNA sequence.

KW TMV-based virus; tobacco mosaic virus; protein isolation; green juice;

KW virus isolation; fusion protein identification; ss.

XX Tobacco mosaic virus.

XX WO9946288-A2.

XX 16-SEP-1999.

XX 09-MAR-1999; 99WO-US05056.

XX 10-MAR-1998; 98US-0037751.

XX (BIOS-) BIOSOURCE TECHNOLOGIES INC.

XX Garger SJ, Holtz RB, McCulloch MJ, Turpen TH;

XX WPI: 1999-561660/47.

XX Obtaining protein, viruses and fusion proteins from plants, using

PT non-denaturing conditions -

XX Disclosure; Page 55-58; 58pp; English.

XX
XX This sequence represents a tobacco mosaic virus (TMV) based virus
CC sequence identified using the method of the invention. The method is for
CC obtaining a soluble protein or peptide of interest from a plant,
CC comprising homogenising the plant to produce green juice, adjusting the
CC pH to less than or equal to 5.2, and heating the juice to a minimum of
CC 45 degrees C. The juice is then centrifuged to produce a supernatant, and
CC the protein or peptide is purified from the supernatant. The method can
CC also be used for obtaining viruses and fusion proteins. The method is
CC especially useful for obtaining IL-1 to IL-10, EPO, G-CSF, GM-CSF,
CC hp-CSF, M-CSF, Factor VIII, Factor IX, tPA, receptors, receptor
CC antagonists, antibodies, single-chain antibodies, enzymes,
CC neuropolypeptides, insulin, antigens, vaccines, peptide hormones,
CC calcitonin, and human growth hormone, or an antimicrobial peptide or
CC protein from protegrins, magainins, cecropins, melittins, indolicidins,
CC defensins, beta-defensins, cryptidins, clavalins, plant defensins,
CC nicin and bactericins, all produced by recombinant means. The new method
CC is more efficient than the prior art for isolating viruses, protein, and
CC peptides. The method is large-scale, and non-denaturing and
CC solvent-limited. Prior art methods do not isolate recombinant proteins,
CC and do not allow fraction 2 proteins to be ultrafiltrated.
XX
XX Sequence 6446 BP; 1873 A; 1234 C; 1563 G; 1776 U; 0 other;

alignment_scores:
Quality: 73.00 Length: 9
Ratio: 8.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-35 x AAZ20646

Align seg 1/1 to: AAZ20646 from: 1 to: 6446

4 LysTrpProTfTrpParGarg 12
|||||
6213 AAGUGGCCUUGUGGCCAUGGCCCGA 6239

seq_name: /SIDS8/gcdata/geneseq/geneseqn/NA2001.DAT:AAF82334

seq_documentation_block:

ID AAF82334 standard; RNA: 6446 BP.

AAF82334;

22-JUN-2001 (first entry)

Tobacco mosaic virus-based coat protein read-through virus TMV661.

TMV; tobacco mosaic virus; TMV661; virus isolation;

non-native protein purification; ribulose 1,5-diphosphate carboxylase;

Rubisco; coat protein read-through; ss.

Tobacco mosaic virus.

WO200119969-A1.

22-MAR-2001.

19-MAY-2000; 2000MO-US13680.

16-SEP-1999; 99US-0397090.

(LARG-) LARGE SCALE BIOLOGY CORP.

Garger SJ, Holtz BR, McCulloch MJ, Turpen TH;

WPI; 2001-328016/34.

Minimizing presence of ribulose 1,5-diphosphate carboxylase to obtain

plant product for isolating bioactive species involves cutting plant

material from plant in cutting period when quantity of Rubisco is at

minimum

Disclosure; Page 72-74; 81pp; English.

The present sequence is a tobacco mosaic virus (TMV)-based virus

which was used to infect field-grown tobacco. The virus was then

isolated from the tobacco plants by a novel process for isolating and

purifying viruses, soluble proteins and peptides from plant sources. In

order to isolate the bioactive species from the undesirable

photosynthetic protein ribulose 1,5-diphosphate carboxylase (Rubisco),

the plant material is cut in a period of the light/dark cycle when the

quantity of Rubisco in the plant is at a minimum. The method is useful

for obtaining a virus of interest. It is also useful for obtaining

soluble recombinant or non-native proteins, such as active mammalian

proteins, enzymes, vaccines, antibodies and peptides, from transgenic

plants.

Sequence 6446 BP; 1873 A; 1234 C; 1563 G; 1776 U; 0 other;

alignment_scores: Quality: 73.00 Length: 9

Ratio: 8.111 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block: US-09-444-281-35 x AAF82334 ..

Align seg 1/1 to: AAF82334 from: 1 to: 6446

4 LysTrpProTfTrpParGarg 12

|||||

6213 AAGUGGCCUUGUGGCCAUGGCCCGA 6239

seq_name: /SIDS8/gcdata/geneseq/geneseqn/NA2000.DAT:AAA28519

seq_documentation_block:
ID AAA28519 standard; DNA: 207 BP.

AAA28519;

29-AUG-2000 (first entry)

PCRIL DNA coding sequence.

Magalain; antimicrobial; transgenic plant; protease degradation; Rev4;

indolicidin; protein production; reverse peptide; ss.

Synthetic.

WO200026344-A1.

11-MAY-2000.

29-OCT-1999; 99WO-US25561.

30-OCT-1998; 98US-0106373.

02-NOV-1998; 98US-0106537.

(INTE-) INTERLINK BIOTECHNOLOGIES LLC.

(KENT) UNIV KENTUCKY RES FOUND.

Everett NP, Li Q, Lawrence C, Davies MH;

WPI; 2000-365597/31.

P-PSDB; AAY92840.

Polypeptides for reducing proteolytic degradation of proteins

administered to, or produced by a plant comprise indolicidin or its

functional equivalents

Example 17; Page 35; 50pp; English.

Indolicidin is a potent antimicrobial tridecapeptide, originally

purified from cytoplasmic granules of bovine neutrophils. Reverse

peptide, Rev4 of indolicidin (see AAY92794) was found to have increased

stability against plant protease degradation. Expression of antimicrobial

peptides in transgenic plants suffers a major limitation in that the

foreign peptides are susceptible to rapid degradation by proteases. The

invention concerns reducing the extent of protease degradation of a

protein applied to, or produced by a plant by administering indolicidin,

Rev4 or a functional equivalent to the plant. Transgenic plants

expressing indolicidin and Rev4 are useful for production of the

antimicrobial peptides. Compositions containing indolicidin and Rev4 are

also useful for production of agronomically important proteins in

plants.

Sequence 207 BP; 49 A; 50 C; 36 G; 72 T; 0 other;

alignment_scores: Quality: 66.00 Length: 10

Ratio: 6.600 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 60.000

alignment_block: US-09-444-281-35 x AAA28519 ..

Align seg 1/1 to: AAA28519 from: 1 to: 207

2 LeuLysLysTrpProTfTrpParGarg 11

|||||

163 ATTAGAGATGCGCTTGCTGCGCTTGGA 192

APPLICANT: Kim, Jeong Hyun
APPLICANT: Hong, Seung-Suh
APPLICANT: Lee, Hyun-Soo
APPLICANT: Samyang Genex Corporation
TITLE OF INVENTION: METHOD FOR MASS PRODUCTION OF
FILE REFERENCE: 6181/0F135
CURRENT APPLICATION NUMBER: US/09/230,180
CURRENT FILING DATE: 1999-03-10
PRIOR APPLICATION NUMBER: PCT/KR98/00132
PRIOR FILING DATE: 1998-05-28
PRIOR APPLICATION NUMBER: KR 13372/1998
PRIOR FILING DATE: 1998-04-09
PRIOR APPLICATION NUMBER: KR 21312/1997
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 36
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 29
LENGTH: 39
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: DNA sequence deduced from Indolicidin peptide
US-09-230-180-29

alignment_scores:
Quality: 73.00 Length: 9
Ratio: 8.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-35 x US-09-230-180-29
Align seg 1/1 to: US-09-230-180-29 from: 1 to: 39

4 LysTrpProTrrPrProTrrParGArg 12
|||||
13 AAATGGCCGTGGTGGCCGTGGCTGCT 39

seq_name: /cgn2_6/ptodata/2/ina/6A_COMB.seq:US-09-259-741-5

seq_documentation_block:

Sequence 5, Application US/09259741
Patent No. 6033895
GENERAL INFORMATION:
APPLICANT: GARGER, STEPHEN
APPLICANT: HOLTZ, R. BARRY
APPLICANT: MCCULLOCH, MICHAEL
APPLICANT: TURPEN, THOMAS
TITLE OF INVENTION: A PROCESS FOR ISOLATING AND
TITLE OF INVENTION: PURIFYING VIRUSES SOLUBLE PROTEINS AND PEPTIDES FROM PLANT
NUMBER OF SEQUENCES: 5
CORRESPONDENCE ADDRESS:
ADDRESSEE: Howrey & Simon
STREET: 1299 Pennsylvania Avenue N.W.
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20004
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/259,741
FILING DATE: February 25, 1999
CLASSIFICATION:
PRIOR APPLICATION DATA:

APPLICATION NUMBER: 09/037,751
FILING DATE: March 10, 1998
ATTORNEY/AGENT INFORMATION:
NAME: Halluin, Albert P
REGISTRATION NUMBER: 25,277
REFERENCE/DOCKET NUMBER: 00801.0140.US01
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-463-8100
TELEFAX: 650-463-8400
TELEX:
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 6446 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: unknown
MOLECULE TYPE: Genomic RNA
US-09-259-741-5

alignment_scores:
Quality: 73.00 Length: 9
Ratio: 8.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-35 x US-09-259-741-5
Align seg 1/1 to: US-09-259-741-5 from: 1 to: 6446

4 LysTrpProTrrPrProTrrParGArg 12
|||||
6213 AAGUGCCUUGUGGCCAAGGCCCGCA 6239

seq_name: /cgn2_6/ptodata/2/ina/6A_COMB.seq:US-09-037-751-5

seq_documentation_block:

Sequence 5, Application US/09037751
Patent No. 6037456
GENERAL INFORMATION:
APPLICANT: GARGER, STEPHEN
APPLICANT: HOLTZ, R. BARRY
APPLICANT: MCCULLOCH, MICHAEL
APPLICANT: TURPEN, THOMAS
TITLE OF INVENTION: A PROCESS FOR ISOLATING AND
TITLE OF INVENTION: PURIFYING VIRUSES SOLUBLE PROTEINS AND PEPTIDES
NUMBER OF SEQUENCES: 5
CORRESPONDENCE ADDRESS:
ADDRESSEE: Howrey & Simon
STREET: 1299 Pennsylvania Avenue N.W.
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20004
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/037,751
FILING DATE: 10-MAR-1998
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Halluin, Albert P
REGISTRATION NUMBER: 25,277
REFERENCE/DOCKET NUMBER: 00801.0140.999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-463-8109

TELEFAX: 650-463-8400
TELEX:
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 6446 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: unknown
MOLECULE TYPE: Genomic RNA
US-09-037-751-5

alignment_scores:
Quality: 73.00 Length: 9
Ratio: 8.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x US-09-037-751-5 ..

Align seg 1/1 to: US-09-037-751-5 from: 1 to: 6446

4 LysTrpProTTrpTTrpTTrpArgArg 12
|||||
6213 AAGUGCCUGGUGGCAUGGCGCGCA 6239

seq_name: /cgn2_6/prodata/2/lna/6B_COMB.seq:US-09-466-422-5

seq_documentation_block:
Sequence 5, Application US/09466422
Patent No. 6303779
GENERAL INFORMATION:
APPLICANT: GARGER, STEPHEN
HOLTZ, R. BARRY
MCCULLOCH, MICHAEL
TURNER, THOMAS
TITLE OF INVENTION: A PROCESS FOR ISOLATING AND
PURIFYING VIRUSES SOLUBLE PROTEINS AND PEPTIDES
*FROM PLANT SOURCES
NUMBER OF SEQUENCES: 5
CORRESPONDENCE ADDRESS:
ADDRESSEE: Howrey & Simon
STREET: 1293 Pennsylvania Avenue N.W.
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20004
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FASTED for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/466,422
FILING DATE: 17-Dec-1999
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 09/037,751
FILING DATE: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: Halluin, Albert P
REGISTRATION NUMBER: 25,277
REFERENCE/DOCKET NUMBER: 00801.0140.999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-463-8109
TELEFAX: 650-463-8400
TELEX: <Unknown>
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 6446 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: unknown

MOLECULE TYPE: Genomic RNA
SEQUENCE DESCRIPTION: SEQ ID NO: 5:
US-09-466-422-5

alignment_scores:
Quality: 73.00 Length: 9
Ratio: 8.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x US-09-466-422-5 ..

Align seg 1/1 to: US-09-466-422-5 from: 1 to: 6446

4 LysTrpProTTrpTTrpTTrpArgArg 12
|||||
6213 AAGUGCCUGGUGGCAUGGCGCGCA 6239

seq_name: /cgn2_6/prodata/2/lna/5A_COMB.seq:US-08-159-784-1

seq_documentation_block:
Sequence 1, Application US/08159784
Patent No. 5643783
GENERAL INFORMATION:
APPLICANT: Bjorn R. Olsen
TITLE OF INVENTION: NOVEL COLLAGEN AND USES THEREOF
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson
STREET: 225 Franklin Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110-2804
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
COMPUTER: IBM PS/2 Model 50z or 55sx
OPERATING SYSTEM: MS-DOS (Version 5.0)
SOFTWARE: WordPerfect (Version 5.1)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/159,784
FILING DATE: December 1, 1993
CLASSIFICATION: 530
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: John F. Freeman
REGISTRATION NUMBER: 29,066
REFERENCE/DOCKET NUMBER: 00246/170001
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 542-5070
TELEFAX: (617) 542-8906
TELEX: 200154
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 4031
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-159-784-1

alignment_scores:
Quality: 60.00 Length: 9
Ratio: 7.500 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 77.778

alignment_block:
US-09-444-281-35 x US-08-159-784-1/rev ..

Align seg 1/1 to reverse of: US-08-159-784-1 from: 1 to: 4031

3 LysLysTrpProTrpTrpProTrp 11
|||||
1671 AATCCTGCTGCTGCTGCTGCTGAGC 1645

seq_name: /cgn2_6/ptodata/2/ina/5B_COMB.seq:US-08-481-337A-1

seq_documentation_block:

Sequence 1, Application US/08481337A
Patent No. 5863738
GENERAL INFORMATION:
APPLICANT: TEN DUKE, Peter
APPLICANT: HELDIN, Carl-Henrik
APPLICANT: MIYAZONO, Kohei
APPLICANT: SAMPATH, Kuber T.
TITLE OF INVENTION: Morphogenic Protein-Specific Cell
TITLE OF INVENTION: Surface Receptors and Uses Therefor
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:
ADDRESSEE: Testa, Hurwitz & Thibault
STREET: 125 High St.
CITY: Boston
STATE: MA
COUNTRY: USA
ZIP: 02110
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/481,337A
FILING DATE: 02-JUN-1995
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: MEYERS, Thomas C.
REGISTRATION NUMBER: 36,989
REFERENCE/DOCKET NUMBER: CRP-097CP2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 248-7000
TELEFAX: (617) 248-7100
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1509 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..1509
OTHER INFORMATION: /product= "Human ALK1"
US-08-481-337A-1

alignment_scores:

Quality:	58.00	Length:	6
Ratio:	9.667	Gaps:	0
Percent Similarity:	100.000	Percent Identity:	100.000

alignment_block:

US-09-444-281-35 x US-08-481-337A-1

Align seg 1/1 to: US-08-481-337A-1 from: 1 to: 1509

5 TrpProTrpTrpProTrp 10
|||||

389 TGCCCTGCTGCTGCTGCTG 406

seq_name: /cgn2_6/ptodata/2/ina/5B_COMB.seq:US-08-696-268B-1

seq_documentation_block:

Sequence 1, Application US/08696268B

Patent No. 5968752

GENERAL INFORMATION:

APPLICANT: ICHIO, Hidenori

APPLICANT: NISHITOH, Hidenori

APPLICANT: SAMPATH, Kuber T.

TITLE OF INVENTION: NOVEL SIGNALING RECEPTOR FOR

NUMBER OF SEQUENCES: 8

CORRESPONDENCE ADDRESS:

ADDRESSEE: Testa, Hurwitz & Thibault

STREET: 125 High St.

CITY: Boston

STATE: MA

COUNTRY: USA

ZIP: 02110

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/696,268B

FILING DATE:

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: MEYERS, Thomas C.

REGISTRATION NUMBER: 36,989

REFERENCE/DOCKET NUMBER: CRP-117

TELECOMMUNICATION INFORMATION:

TELEPHONE: (617) 248-7000

TELEFAX: (617) 248-7100

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 1509 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: cDNA

FEATURE:

NAME/KEY: CDS

LOCATION: 1..1509

OTHER INFORMATION: /product= "Human ALK-1"

US-08-696-268B-1

alignment_scores:

Quality:	58.00	Length:	6
Ratio:	9.667	Gaps:	0
Percent Similarity:	100.000	Percent Identity:	100.000

alignment_block:

US-09-444-281-35 x US-08-696-268B-1

Align seg 1/1 to: US-08-696-268B-1 from: 1 to: 1509

5 TrpProTrpTrpProTrp 10
|||||

389 TGCCCTGCTGCTGCTGCTG 406

seq_name: /cgn2_6/ptodata/2/ina/PCTUS_COMB.seq:PCT-US95-05467-1

seq_documentation_block:

Sequence 1, Application PC/TUS9505467

GENERAL INFORMATION:

APPLICANT:

APPLICANT:

TITLE OF INVENTION: MORPHOGENIC PROTEIN-SPECIFIC CELL

TITLE OF INVENTION: SURFACE RECEPTORS AND USES THEREFOR

NUMBER OF SEQUENCES: 15

CORRESPONDENCE ADDRESS:

ADDRESSEE: PATENT ADMINISTRATOR, TESTA, HURWITZ &

STREET: 53 STATE STREET

CITY: BOSTON
STATE: MA
COUNTRY: USA
ZIP: 02109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/05467
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: PITCHER, EDMUND R.
REGISTRATION NUMBER: 27,829
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 248-7000
TELEFAX: (617) 248-7100
SEQUENCE CHARACTERISTICS:
LENGTH: 1509 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..1509
OTHER INFORMATION: /product= "Human ALK1"
PCT-US95-05467-1

alignment_scores:
Quality: 58.00 Length: 6
Ratio: 9.667 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x PCT-US95-05467-1 ..

Align seg 1/1 to: PCT-US95-05467-1 from: 1 to: 1509

5 TrpProrTrrProrTrr 10
|||||
389 TGCCCTGTGTCCTGG 406

seq_name: /cgn2_6/prodata/2/ina/PCTUS_COMB.seq:PCT-US94-11328A-3

seq_documentation_block:

Sequence 3, Application PC/TUS9411328A
GENERAL INFORMATION:
APPLICANT: HE, ET AL.
TITLE OF INVENTION: TAR-1 and TAR-3
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: CARELLA, HYRNE, BAIN, GILFILLAN,
CECCHI, STEWART & OLSTEIN
STREET: 6 BECKER FARM ROAD
CITY: ROSELAND
STATE: NEW JERSEY
COUNTRY: USA
ZIP: 07068
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5 INCH DISKETTE
COMPUTER: IBM PS/2
OPERATING SYSTEM: MS-DOS
SOFTWARE: WORD PERFECT 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US94/11328A
FILING DATE: Submitted herewith
CLASSIFICATION:

PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: FERRARO, GREGORY D.
REGISTRATION NUMBER: 36,134
REFERENCE/DOCKET NUMBER: 325800-132
TELECOMMUNICATION INFORMATION:
TELEPHONE: 201-994-1700
TELEFAX: 201-994-1744
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 1596 BASE PAIRS
TYPE: NUCLEIC ACID
STRANDEDNESS: SINGLE
TOPOLOGY: LINEAR
MOLECULE TYPE: CDNA
PCT-US94-11328A-3

alignment_scores:
Quality: 58.00 Length: 6
Ratio: 9.667 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x PCT-US94-11328A-3 ..

Align seg 1/1 to: PCT-US94-11328A-3 from: 1 to: 1596

5 TrpProrTrrProrTrr 10
|||||
470 TGCCCTGTGTCCTGG 487

seq_name: /cgn2_6/prodata/2/ina/GB_COMB.seq:US-09-382-256-1

seq_documentation_block:

Sequence 1, Application US/09382256A
Patent No. 6207814
GENERAL INFORMATION:
APPLICANT: MIYAZONO, Kohel
TEN DIJKE, Peter
FRANZEN, Petra
YAMASHITA, Hidetoshi
HELDIN, Carl-Heinrik
TITLE OF INVENTION: ACTIVIN RECEPTOR LIKE KINASES, PROTEINS
HAVING SERINE THREONINE KINASE DOMAINS,
AND THEIR USE
NUMBER OF SEQUENCES: 29
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pulbright & Jaworski L.L.P.
STREET: 666 Fifth Avenue
CITY: New York City
STATE: New York
COUNTRY: USA
ZIP: 10103
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.25 inch, 1.44mb
COMPUTER: IBM PS/2
OPERATING SYSTEM: PC-DOS
SOFTWARE: Wordperfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/382,256A
FILING DATE: 24-Aug-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/GB93/02367
FILING DATE: No. 6207814ember 17, 1993
APPLICATION NUMBER: GB 9224057.1
FILING DATE: No. 6207814ember 17, 1992
APPLICATION NUMBER: GB 9304677.9
FILING DATE: March 8, 1993
APPLICATION NUMBER: GB 9304680.3

```

? FILING DATE: March 8, 1993
? APPLICATION NUMBER: 9311047.6
? FILING DATE: May 28, 1993
? APPLICATION NUMBER: 9313763.6
? FILING DATE: July 2, 1993
? APPLICATION NUMBER: 9316099.2
? FILING DATE: August 3, 1993
? APPLICATION NUMBER: 321344.5
? FILING DATE: October 15, 1993
? ATTORNEY/AGENT INFORMATION:
?   NAME: No. 6207814man D. Hanson
?   REGISTRATION NUMBER: 30,946
? TELECOMMUNICATION INFORMATION:
?   TELEPHONE: (212) 318-3000
?   TELEFAX: (212) 752-5958
? INFORMATION FOR SEQ ID NO: 1:
?   LENGTH: 1984 base pairs
?   TYPE: nucleic acid
?   STRANDEDNESS: unknown
?   TOPOLOGY: linear
?   MOLECULE TYPE: CDNA
?   HYPOTHETICAL: NO
?   ANTI-SENSE: NO
?   FRAGMENT TYPE: Internal
?   ORIGINAL SOURCE:
?     ORGANISM: Homo sapiens
?   FEATURE:
?     NAME/KEY: CDS
?     LOCATION: 283..1791
?   SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-382-256-1

alignment_scores:
?   Quality: 58.00      Length: 6
?   Ratio: 9.667      Gaps: 0
?   Percent Similarity: 100.000      Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x US-09-382-256-1 ..
Align seg 1/1 to: US-09-382-256-1 from: 1 to: 1984
5 TrpProtTrpProtTrp 10
|||||
671 TGGCCCTGTGTGGCCCTGG 688

seq_name: /cgn2_6/ptodata/2/ina/6A_COMB.seq:US-09-395-115-1
seq_documentation_block:
? Sequence 1, Application US/09395115
? Patent No. 6271365
? GENERAL INFORMATION:
?   APPLICANT: Miyazono, Kohel; DiJke, Peter Ten;
?   TITLE OF INVENTION: Activin Receptor-Like Kinase, Proteins
?   TITLE OF INVENTION: Having Serine Threonine Kinase Domains And Their Use
?   NUMBER OF SEQUENCES: 29
?   CORRESPONDENCE ADDRESS:
?     ADDRESSEE: Felfe & Lynch
?     STREET: 805 Third Avenue
?     CITY: New York City
?     STATE: New York
?     ZIP: 10022
? COMPUTER READABLE FORM:
?   MEDIUM TYPE: Diskette, 3.5 inch, 360 kb storage
?   COMPUTER: IBM
?   OPERATING SYSTEM: PC-DOS
?   SOFTWARE: Wordperfect
?   CURRENT APPLICATION DATA:
?     APPLICATION NUMBER: US/09/395,115
```

```

? FILING DATE:
? CLASSIFICATION:
? PRIOR APPLICATION DATA:
?   APPLICATION NUMBER: US/08/436,265
?   FILING DATE: 30-October-1995
?   APPLICATION NUMBER: PCT/GB93/02367
?   FILING DATE: 17-No. 6271365ember-1993
?   PRIOR APPLICATION DATA:
?     APPLICATION NUMBER: 9224057.1
?     FILING DATE: 17-No. 6271365ember-1992
?   PRIOR APPLICATION DATA:
?     APPLICATION NUMBER: 9304677.9
?     FILING DATE: 8-March-1993
?   PRIOR APPLICATION DATA:
?     APPLICATION NUMBER: 9304680.3
?     FILING DATE: 8-March-1993
?   PRIOR APPLICATION DATA:
?     APPLICATION NUMBER: 9311047.6
?     FILING DATE: 28-May-1993
?   PRIOR APPLICATION DATA:
?     APPLICATION NUMBER: 9313763.6
?     FILING DATE: 2-July-1993
?   PRIOR APPLICATION DATA:
?     APPLICATION NUMBER: 9136099.2
?     FILING DATE: 3-August-1993
?   PRIOR APPLICATION DATA:
?     APPLICATION NUMBER: 9321344.5
?     FILING DATE: 15-October-1993
?   ATTORNEY/AGENT INFORMATION:
?     NAME: Kohlei, Vineet
?     REGISTRATION NUMBER: 37,003
?     REFERENCE/DOCKET NUMBER: LUD 5298
?   TELECOMMUNICATION INFORMATION:
?     TELEPHONE: (212) 688-9200
?     TELEFAX: (212) 838-3884
?   INFORMATION FOR SEQ ID NO: 1:
?     SEQUENCE CHARACTERISTICS:
?       LENGTH: 1984 base pairs
?       TYPE: nucleic acid
?       STRANDEDNESS: unknown
?       TOPOLOGY: linear
?       MOLECULE TYPE: CDNA
?       HYPOTHETICAL: NO
?       ANTI-SENSE: NO
?       FRAGMENT TYPE: Internal
?       ORIGINAL SOURCE:
?         ORGANISM: Homo sapiens
?       FEATURE:
?         NAME/KEY: CDS
?         LOCATION: 283..1791
?     US-09-395-115-1

alignment_scores:
?   Quality: 58.00      Length: 6
?   Ratio: 9.667      Gaps: 0
?   Percent Similarity: 100.000      Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x US-09-395-115-1 ..
Align seg 1/1 to: US-09-395-115-1 from: 1 to: 1984
5 TrpProtTrpProtTrp 10
|||||
671 TGGCCCTGTGTGGCCCTGG 688

seq_name: /cgn2_6/ptodata/2/ina/6A_COMB.seq:US-08-749-816-1
seq_documentation_block:
? Sequence 1, Application US/08749816
? Patent No. 6013470
? GENERAL INFORMATION:
```

APPLICANT: Lesage, Florian
APPLICANT: Guillemae, Eric
APPLICANT: Fink, Michel
APPLICANT: Duprat, Fabrice
APPLICANT: Lazdunski, Michel
APPLICANT: Romey, Georges
APPLICANT: Barhanin, Jacques
TITLE OF INVENTION: FAMILY OF MAMMALIAN POTASSIUM CHANNELS,
TITLE OF INVENTION: THEIR CLONING AND THEIR USE ESPECIALLY FOR THE SCREENING
OF DRUGS
NUMBER OF SEQUENCES: 19
TITLE OF INVENTION: OF DRUGS
CORRESPONDENCE ADDRESS:
ADDRESSEE: WEISER & ASSOCIATES
STREET: 230 South Fifteenth Street, Suite 500
City: Philadelphia
STATE: PA
COUNTRY: USA
ZIP: 19102
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/749,816
FILING DATE: 15-NOV-1996
CLASSIFICATION: 536
ATTORNEY/AGENT INFORMATION:
NAME: Weiser, Gerard J.
REGISTRATION NUMBER: 19,763
REFERENCE/DOCKET NUMBER: 989,635IP
TELECOMMUNICATION INFORMATION:
TELEPHONE: 215-875-8883
TELEFAX: 215-875-8394
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1894 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 183..1190
US-08-749-816-1
alignment_scores:
Quality: 57.50 Length: 13
Ratio: 5.227 Gaps: 1
Percent Similarity: 84.615 Percent Identity: 61.538
alignment_block:
US-09-444-281-35 x US-08-749-816-1
Align seg 1/1 to: US-08-749-816-1 from: 1 to: 1894
2 LeuysysTrpProTrp...TrrProtrparGaglyS 13
|||||:|||||:||||| 111 |||||:|||||:
136 CTGCGCGCTTGCGCTTGGCTTGGCGCGCGCGG 174
seq_name: /cgn2_6/plodata//lha/58_COMB.seq:US-08-149-097D-24
seq_documentation_block:
Sequence 24, Application US/08149097D
GENERAL INFORMATION:
APPLICANT: Harpold, Michael
APPLICANT: Ellis, Steven
APPLICANT: Williams, Mark
APPLICANT: Feldman, Daniel
APPLICANT: McCue, Ann
APPLICANT: Brenner, Robert

TITLE OF INVENTION: HUMAN CALCIUM CHANNEL COMPOSITIONS AND
METHODS
TITLE OF INVENTION: METHODS
NUMBER OF SEQUENCES: 40
CORRESPONDENCE ADDRESS:
ADDRESSEE: Brown, Martin, Haller & McClain
STREET: 1660 Union Street
City: San Diego
STATE: California
COUNTRY: USA
ZIP: 92101-2926
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/149,097D
FILING DATE: 05-NOV-1993
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/105,536
FILING DATE: 11-AUG-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO PCT/US92/06903
FILING DATE: 14-AUG-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/914,231
FILING DATE: 13-JUL-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/868,354
FILING DATE: 10-APR-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/745,206
FILING DATE: 15-AUG-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/620,250
FILING DATE: 30-NOV-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/482,384
FILING DATE: 20-FEB-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/603,751
FILING DATE: 04-APR-1989
PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO PCT/US89/01408
FILING DATE: 04-APR-1989
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/176,899
FILING DATE: 04-APR-1988
ATTORNEY/AGENT INFORMATION:
NAME: Seidman, Stephanie L.
REGISTRATION NUMBER: 33,779
REFERENCE/DOCKET NUMBER: 6362-55038
TELECOMMUNICATION INFORMATION:
TELEPHONE: (619) 238-0062
TELEFAX: (619) 238-0062
INFORMATION FOR SEQ ID NO: 24:
SEQUENCE CHARACTERISTICS:
LENGTH: 7032 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: 169..6921
OTHER INFORMATION: /product="Alphale-1 subunit of
human calcium channel"
US-08-149-097D-24
alignment_scores:
Quality: 57.50 Length: 11

Ratio: 6.389 Gaps: 1
Percent Similarity: 81.818 Percent Identity: 72.727

alignment_block:
US-09-444-281-35 x US-08-149-097D-24 ...

Align seg 1/1 to: US-08-149-097D-24 from: 1 to: 7032

2 LeuLysLysTrpProTrp...TrpProTrpArg 11
||||| ||||||| ||||||| |||||||
2606 TTGAGGGCCTGGCCCTGGCCCTGGCCCTGGAGA 2638

seq_name: /cgn2.6/ptodata/2/ina/6A.COMB.seq:US-08-949-386-24

seq_documentation_block:
Sequence 24, Application US/08949386

Patent No. 6090623

GENERAL INFORMATION:
APPLICANT: Harpold, Michael

APPLICANT: Ellis, Steven

APPLICANT: Williams, Mark

APPLICANT: McCue, Ann

APPLICANT: Gillespie, Allison

TITLE OF INVENTION: HUMAN CALCIUM CHANNEL COMPOSITIONS AND

NUMBER OF SEQUENCES: 38

CORRESPONDENCE ADDRESS:

ADDRESSEE: Brown, Martin, Haller & McClain

STREET: 1660 Union Street

CITY: San Diego

STATE: California

COUNTRY: US

ZIP: 92101

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patent Release #1.0, Version #1.25

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/949,386

FILING DATE:

CLASSIFICATION:

Prior Application DATA:
APPLICATION NUMBER: US/08/290,012

FILING DATE: 11-AUG-1994

APPLICATION NUMBER: 08/149,097

FILING DATE: 5-NOV-1993

Prior Application DATA:
APPLICATION NUMBER: 08/105,536

FILING DATE: 11-AUG-1993

ATTORNEY/AGENT INFORMATION:
NAME: Seidman, Stephanie L.

REGISTRATION NUMBER: 33,779

REFERENCE/DOCKET NUMBER: 519808

TELECOMMUNICATION INFORMATION:
TELEPHONE: (619) 238-0999

TELEFAX: (619) 238-0062

INFORMATION FOR SEQ ID NO: 24:

SEQUENCE CHARACTERISTICS:
LENGTH: 7032 base pairs

TYPE: nucleic acid

STRANDEDNESS: double

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

FEATURE:

NAME/KEY: CDS

LOCATION: 166..6921

OTHER INFORMATION: /standard_name="Alpha-1E-1"

US-08-949-386-24

Ratio: 6.389 Gaps: 1
Percent Similarity: 81.818 Percent Identity: 72.727

alignment_block:
US-09-444-281-35 x US-08-949-386-24 ..

Align seg 1/1 to: US-08-949-386-24 from: 1 to: 7032

2 LeuLysLysTrpProTrp...TrpProTrpArg 11
||||| ||||||| ||||||| |||||||
2606 TTGAGGGCCTGGCCCTGGCCCTGGCCCTGGAGA 2638

alignment_scores: Quality: 57.50 Length: 11

```

/SIDS8/gcgdata/geneseq/geneseqn/NA2001.DAT:AAF83651 + 39.00 98.94 2.0e+03 5523
/SIDS8/gcgdata/geneseq/geneseqn/NA2001.DAT:AAF24892 + 39.00 87.83 8.1e+03 2038
/SIDS8/gcgdata/geneseq/geneseqn/NA2000.DAT:AAF11976 + 38.00 114.22 276.00 629
/SIDS8/gcgdata/geneseq/geneseqn/NA1994.DAT:AAQ54498 + 38.00 112.49 344.58 771
/SIDS8/gcgdata/geneseq/geneseqn/NA1995.DAT:AAQ02565 + 38.00 112.49 344.58 771

seq_name: /SIDS8/gcgdata/geneseq/geneseqn/NA2000.DAT:AAA27297
seq_documentation_block:
ID AAA27297 standard; DNA; 97 BP.
XX
XX AAA27297;
AC
XX
XX
XX 20-SEP-2000 (first entry)
XX
XX Oligonucleotide used for synthesis of anionic spacer cassette.
DE
XX Oligonucleotide; cellulose binding domain; CBD; cationic peptide;
KW MB1-11; indolicidin; bovine; ss.
XX
XX Synthetic.
OS
XX WO200031279-A2.
XX
XX 02-JUN-2000.
XX
XX 19-NOV-1999; 99WO-CA01107.
XX
XX 20-NOV-1998; 98US-0109218.
XX
XX (MICR-) MICROLOGIX BIOTECH INC.
XX
XX Burian J, Bartfeld D;
XX
XX WPI; 2000-400086/34.
XX
XX Multi-domain fusion protein expression cassette used for high yield
PT stable production of foreign peptide gene products -
PT
XX
XX Example 5; Page 40; 73pp; English.
XX
XX A novel method allows the efficient production of cationic peptides in
CC recombinant host cells. The method involves construction of a
CC multi-domain fusion protein expression cassette comprising a promoter and
CC a nucleic acid molecule expressed as an insoluble protein. The inclusion
CC of anionic peptide sequences in the linker sequences neutralises the
CC positive charge of the cationic peptide so that the charge of the
CC fusion protein is controlled. This cassette allows high yield, stable
CC production of the cationic peptide. Cationic peptides such as
CC bovine indolicidin may be used as antimicrobial agents. The present
CC sequence is an oligonucleotide that was used to synthesise the
CC anionic spacer of the expression cassette.
XX
XX
XX Sequence 97 BP; 23 A; 29 C; 26 G; 19 T; 0 other;
SQ

alignment_scores:
Quality: 54.00 Length: 10
Ratio: 5.400 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-27 x AAA27297 ..

Align seg 1/1 to: AAA27297 from: 1 to: 97

1 HHSGLUALGLUPROGLUALGLUPROILE 10
|||||
38 CATGAAGCGAGACCGAAGCGAACCAGATT 67

seq_name: /SIDS8/gcgdata/geneseq/geneseqn/NA1999.DAT:AAV90541
seq_documentation_block:

```

```

ID  AAV90541 standard; cDNA; 375 BP.
XX
AC  AAV90541;
XX
DE  15-FEB-1999 (first entry)
XX
DE  EST clone BK517.
XX
KM  Human; secreted protein; expressed sequence tag; EST; haematopoiesis;
KM  tissue growth; activin; inhibin; chemotaxis; chemokinesis; haemostatic;
KM  receptor; ligand; thrombolytic; anti-inflammatory; cadherin; anti-tumour;
KM  gene therapy; ss.
XX
OS  Homo sapiens.
XX
PN  M09845436-A2.
XX
PD  15-OCT-1998.
XX
PF  10-APR-1998; 98MO-US06955.
XX
PR  10-APR-1997; 97US-0838821.
XX
PA  (GEM ) GENETICS INST INC.
XX
PI  Agostino MJ, Jacobs K, Lavallie ER, McCoy JM, Merberg D;
PI  Racie LA, Spaulding V, Treacy M;
XX
DR  MPI; 1999-070077/06.
XX
PT  New polynucleotides encoding human secreted proteins - derived from
PT  e.g. human blood, kidney, foetal lung, placenta, testes, brain,
PT  ovary, pituitary, retina and colon cDNA libraries.
XX
PS  Claim 1; Page 574; 618p; English.
XX
CC  The present sequence represents a human expressed sequence tag (EST).
CC  The polynucleotide, which is a secreted EST, and the encoded protein
CC  are predicted to have useful biological activities which would make
CC  them suitable for treating, preventing or ameliorating medical
CC  conditions in humans and animals, although no supporting data is
CC  given. Suggested activities include nutritional activity, immune
CC  stimulating or suppressing activity, haematopoiesis regulating
CC  activity, tissue growth activity, activin/inhibin activity,
CC  chemotactic/chemokinetic activity, haemostatic and thrombolytic
CC  activity, receptor/ligand activity, anti-inflammatory activity,
CC  cadherin/tumour invasion suppressor activity, tumour inhibition
CC  activity. The polynucleotide may also be useful for gene therapy.
XX
SQ  Sequence 375 BP; 56 A; 125 C; 113 G; 81 T; 0 other;

alignment_scores:
    Quality: 41.00      Length: 9
    Ratio: 4.556        Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:
US-09-444-281-27 x AAV90541/rev ..

Align seg 1/1 to reverse of: AAV90541 from: 1 to: 375

1 HisGluAlaGluProGluAlaGluPro 9
|||||
140 CACGAAGCAGAGCCTGAGGGAGCCCG 114

seq_name: /SID8/gcdata/geneseq/geneseq/NA2001.DAT:AAH10202
seq_documentation_block:
ID  AAH10202 standard; cDNA; 557 BP.
XX
AC  AAH10202;
XX

```

```

DT  26-JUN-2001 (first entry)
XX
DE  Human cDNA clone (3'-primer) SEQ ID NO:7037.
XX
DE  Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.
XX
OS  Homo sapiens.
XX
PN  EP1074617-A2.
XX
PD  07-FEB-2001.
XX
PF  28-JUL-2000; 2000EP-0116126.
XX
PR  29-JUL-1999; 99JP-0248036.
PR  27-AUG-1999; 99JP-0300253.
PR  11-JAN-2000; 2000JP-0118776.
PR  02-MAY-2000; 2000JP-0183767.
PR  09-JUN-2000; 2000JP-0241899.
XX
PA  (HELI-) HELIX RES INST.
XX
PI  Ota T, Isegai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PI  Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX
DR  MPI; 2001-318749/34.
XX
PT  Primer sets for synthesizing polynucleotides, particularly the 5602
PT  full-length cDNAs defined in the specification, and for the detection
PT  and/or diagnosis of the abnormality of the proteins encoded by the
PT  full-length cDNAs -
XX
PS  Claim 3; SEQ ID 7037; 2537pp + CD ROM; English.
XX
CC  The present invention describes primer sets for synthesizing 5602
CC  full-length cDNAs defined in the specification. Where a primer set
CC  comprises: (a) an oligo-dr primer and an oligonucleotide complementary
CC  to the complementary strand of a polynucleotide which comprises one of
CC  the 5602 nucleotide sequences defined in the specification, where the
CC  oligonucleotide comprises at least 15 nucleotides; or (b) a combination
CC  of an oligonucleotide comprising a sequence complementary to the
CC  complementary strand of a polynucleotide which comprises a 5'-end
CC  sequence and an oligonucleotide comprising a sequence complementary to a
CC  polynucleotide which comprises a 3'-end sequence, where the
CC  oligonucleotide comprises at least 15 nucleotides and the combination of
CC  the 5'-end sequence/3'-end sequence is selected from those defined in
CC  the specification. The primer sets can be used in antisense therapy and
CC  in gene therapy. The primers are useful for synthesizing polynucleotides,
CC  particularly full-length cDNAs. The primers are also useful for the
CC  detection and/or diagnosis of the abnormality of the proteins encoded by
CC  the full-length cDNAs. The primers allow obtaining of the full-length
CC  cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
CC  AAH13633 to AAH18742 represent human cDNA sequences; AAH92446 to
CC  AAH95893 represent human amino acid sequences; and AAH13629 to AAH13632
CC  represent oligonucleotides, all of which are used in the exemplification
CC  of the present invention.
XX
SQ  Sequence 557 BP; 99 A; 165 C; 183 G; 103 T; 7 other;

alignment_scores:
    Quality: 41.00      Length: 9
    Ratio: 4.556        Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:
US-09-444-281-27 x AAH10202 ..

Align seg 1/1 to: AAH10202 from: 1 to: 557

1 HisGluAlaGluProGluAlaGluPro 9
|||||
253 CACGAAGCAGAGCCTGAGGGAGCCCG 279

```

```

seq_name: /SID58/gcgdata/geneseq/geneseqn/NA1991.DAT:AAQ12864
seq_documentation_block:
ID   AAQ12864 standard; DNA: 849 BP.
XX
XX   AAQ12864;
AC
XX   11-OCT-1991 (first entry)
DT
XX   Human Cytotoxic Cell Protease-1 coding sequence.
DE
XX   hccp1 inhibitor; cytotoxic T-lymphocytes; ss.
KM
XX   Homo sapiens.
OS
XX   WO9110685-A.
PN
XX   25-JUL-1991.
PD
XX   17-JAN-1991; 91WO-US00340.
PF
XX   19-JAN-1990; 90US-0467880.
PR
XX   (SERA-) SERAGEN INC.
PA
XX   Bleackley RC, Lobe CG, Paetkau VH, James MN, Murphy M;
PI
XX   WPI; 1991-237989/32.
DR
XX
XX   DNA vectors, and inhibitors of cytotoxic cell protease - for
PT
XX   treatment of auto-immune diseases e.g. pernicious anaemia,
PT
XX   rheumatoid arthritis, allo-graft rejection etc.
PS
XX   Disclosure; Fig 6; 62pp; English.
XX
XX   The hccp1 coding sequence was isolated from cytotoxic T-cell
CC
XX   lymphocytes. Vectors comprising the hccp1 coding sequence are
CC
XX   claimed. Clone hcc1 was isolated and found to be the human analogue
CC
XX   of murine C11.
CC
XX   See AAQ12862-6 and AAR13254-R13262.
CC
XX
XX   Sequence 849 BP; 232 A; 230 C; 222 G; 165 T; 0 other;
SQ
alignment_scores:
Quality: 41.00 Length: 11
Ratio: 4.100 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 63.636
alignment_block:
US-09-444-281-27 x AAQ12864 ..
Align seg 1/1 to: AAQ12864 from: 1 to: 849
1 HisGLuAlaGluProGluAlaGluProIleMet 11
|||||
68 CATGAGGCCGAGCCCACTCCGCCCTACATG 100
seq_name: /SID58/gcgdata/geneseq/geneseqn/NA2000.DAT:AAA28524
seq_documentation_block:
ID   AAA28524 standard; cDNA: 1b01 BP.
XX
XX   AAA28524;
AC
XX   29-AUG-2000 (first entry)
DT
XX   Human oploïd growth factor receptor cDNA spliced version 1.
DE
XX
XX   OGFR: oploïd growth factor receptor; growth inhibitor; proliferative;
KM
XX   cytostatic; vulnerrary. gene therapy; antagonist; chromosome 20q13.3; ss.
XX

```

```

OS   Homo sapiens.
XX
XX   Key
FH   5'UTR
FT
XX   CDS
FT
XX   3'UTR
FT
XX
XX   WO200026340-A2.
PN
XX   11-MAY-2000.
PD
XX   02-NOV-1999; 99WO-US25802.
PE
XX   03-NOV-1998; 98US-0106879.
PR
XX   (PENN-) PENN STATE RES FOUND.
PA
XX   Zagon IS, McLaughlin PJ, Verderame MF;
PI
XX   WPI; 2000-365594/31.
DR
XX   P-PSDB; AAY92810.
DR
XX
XX   New cDNA encoding rat and human oploïd growth factor receptors which
PT
XX   modulate cell growth, useful for treating cancer
PT
XX   Claim 1; Page 82-83; 91pp; English.
PS
XX
XX   Primers generated from rat oploïd growth factor receptor (OGFR) cDNA were
CC
XX   used to clone a fragment of the human OGFR cDNA. The complete sequence of
CC
XX   human OGFR was assembled with a combination of 3' and 5' RACE. 5' RACE
CC
XX   consistently yielded a single species of cDNA, while the 3' RACE revealed
CC
XX   extensive alternative splicing. The alternate splice forms were missing
CC
XX   the imperfect repeats or differed in the number of imperfect repeats. The
CC
XX   human OGFR gene chromosomal location was determined by FISH as 20q13.3.
CC
XX   OGFR proteins, nucleic acid molecules, antibodies, transformed cells and
CC
XX   expression vector are useful for detecting expression or levels of an
CC
XX   OGFR in a tissue. OGFR nucleic acids can be used to inhibit growth of
CC
XX   cells in vitro. The antisense sequences and antibodies can be used to
CC
XX   promote growth of cells in vitro. Cell growth can be promoted by
CC
XX   interfering with the OGFR ligand-receptor system, especially where a
CC
XX   subject suffers from a tissue wound. Treating cancer comprises enhancing
CC
XX   the function of the OGFR ligand-receptor system in cancerous cells of a
CC
XX   patient or administering the OGFR nucleic acid to the patient.
SQ
Sequence 1601 BP; 322 A; 485 C; 558 G; 236 T; 0 other;
alignment_scores:
Quality: 41.00 Length: 9
Ratio: 4.556 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778
alignment_block:
US-09-444-281-27 x AAA28524/rev ..
Align seg 1/1 to reverse of: AAA28524 from: 1 to: 1601
1 HisGLuAlaGluProGluAlaGluPro 9
|||||
1418 CACGAAGCAGAGCTGAGGAGGCCG 1392
seq_name: /SID58/gcgdata/geneseq/geneseqn/NA2000.DAT:AAA28526
seq_documentation_block:
ID   AAA28526 standard; cDNA: 2289 BP.
XX
XX   AAA28526;
AC
XX   29-AUG-2000 (first entry)
DT

```

```
XX DE Human opioid growth factor receptor cDNA spliced version 7.
XX AC
XX KM OGRF: opioid growth factor receptor; growth inhibitor; proliferative;
XX KM cytostatic; vulnerrary. gene therapy; antagonist; chromosome 20q13.3; ss.
XX OS Homo sapiens.
XX FH Key
XX FT 5'UTR 1..33 Location/Qualifiers
XX FT CDS /tag= a
XX FT /tag= b
XX FT /product= Opioid_growth_factor_receptor
XX FT 3'UTR 2008..2289
XX FT /tag= c
XX PN WO200026340-A2.
XX PD 11-MAY-2000.
XX PE 02-NOV-1999; 99WO-US25802.
XX PR 03-NOV-1998; 98US-0106879.
XX PA (PENN-) PENN STATE RES FOUND.
XX PI Zagon IS, McLaughlin PJ, Verderame MF;
XX DR MPI: 2000-365594/31.
XX DR P-PSDB; AAY92812.
XX PT New cDNA encoding rat and human opioid growth factor receptors which
XX PT modulate cell growth, useful for treating cancer
XX PS Claim 1; Page 87-89; 91pp; English.
XX PS
XX CC Primers generated from rat opioid growth factor receptor (OGFR) cDNA were
XX CC used to clone a fragment of the human OGFR cDNA. The complete sequence of
XX CC human OGFR was assembled with a combination of 3' and 5' RACE. 5' RACE
XX CC consistently yielded a single species of cDNA, while the 3' RACE revealed
XX CC extensive alternative splicing. The alternate splice forms were missing
XX CC the imperfect repeats or differed in the number of imperfect repeats. The
XX CC human OGFR gene chromosomal location was determined by FISH as 20q13.3.
XX CC OGFR proteins, nucleic acid molecules, antibodies, transformed cells and
XX CC expression vector are useful for detecting expression or levels of an
XX CC OGFR in a tissue. OGFR nucleic acids can be used to inhibit growth of
XX CC cells in vitro. The antisense sequences and antibodies can be used to
XX CC promote growth of cells in vitro. Cell growth can be promoted by
XX CC interfering with the OGR ligand-receptor system, especially where a
XX CC subject suffers from a tissue wound. Treating cancer comprises enhancing
XX CC the function of the OGR ligand-receptor system in cancerous cells of a
XX CC patient or administering the OGFR nucleic acid to the patient.
XX SQ Sequence 2289 BP; 470 A; 714 C; 809 G; 296 T; 0 other;

alignment_scores:
    Quality: 41.00 Length: 9
    Ratio: 4.556 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:
US-09-444-281-27 x AAA28526/rev ..
Align seg 1/1 to reverse of: AAA28526 from: 1 to: 2289
1 HHSGLUAlaGluProGluAlaGluPro 9
|||||
2105 CACGAAGCAGAGCCTGAGGCGAGCCCG 2079
seq_name: /SID58/gcdata/geneseq/geneseqn/NA2000.DAT:AAA28522
```

```
seq_documentation_block:
ID AAA28522 standard; cDNA: 2290 BP.
XX AC
XX AC AAA28522;
XX DT 29-AUG-2000 (first entry)
XX DE Human opioid growth factor receptor cDNA of spliced form A.
XX KM OGRF: opioid growth factor receptor; growth inhibitor; proliferative;
XX KM cytostatic; vulnerrary. gene therapy; antagonist; chromosome 20q13.3; ss.
XX OS Homo sapiens.
XX FH Key
XX FT CDS 34..2007 Location/Qualifiers
XX FT /tag= a
XX PN WO200026340-A2.
XX PD 11-MAY-2000.
XX PE 02-NOV-1999; 99WO-US25802.
XX PR 03-NOV-1998; 98US-0106879.
XX PA (PENN-) PENN STATE RES FOUND.
XX PI Zagon IS, McLaughlin PJ, Verderame MF;
XX DR MPI: 2000-365594/31.
XX PT New cDNA encoding rat and human opioid growth factor receptors which
XX PT modulate cell growth, useful for treating cancer
XX PS Claim 1; Page 77-78; 91pp; English.
XX PS
XX CC Primers generated from rat opioid growth factor receptor (OGFR) cDNA were
XX CC used to clone a fragment of the human OGFR cDNA. The complete sequence of
XX CC human OGFR was assembled with a combination of 3' and 5' RACE. 5' RACE
XX CC consistently yielded a single species of cDNA, while the 3' RACE revealed
XX CC extensive alternative splicing. The alternate splice forms were missing
XX CC the imperfect repeats or differed in the number of imperfect repeats. The
XX CC human OGFR gene chromosomal location was determined by FISH as 20q13.3.
XX CC OGFR proteins, nucleic acid molecules, antibodies, transformed cells and
XX CC expression vector are useful for detecting expression or levels of an
XX CC OGFR in a tissue. OGFR nucleic acids can be used to inhibit growth of
XX CC cells in vitro. The antisense sequences and antibodies can be used to
XX CC promote growth of cells in vitro. Cell growth can be promoted by
XX CC interfering with the OGR ligand-receptor system, especially where a
XX CC subject suffers from a tissue wound. Treating cancer comprises enhancing
XX CC the function of the OGR ligand-receptor system in cancerous cells of a
XX CC patient or administering the OGFR nucleic acid to the patient.
XX SQ Sequence 2290 BP; 470 A; 713 C; 807 G; 297 T; 3 other;

alignment_scores:
    Quality: 41.00 Length: 9
    Ratio: 4.556 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:
US-09-444-281-27 x AAA28522/rev ..
Align seg 1/1 to reverse of: AAA28522 from: 1 to: 2290
1 HHSGLUAlaGluProGluAlaGluPro 9
|||||
2106 CACGAAGCAGAGCCTGAGGCGAGCCCG 2080
seq_name: /SID58/gcdata/geneseq/geneseqn/NA2000.DAT:AAA28525
seq_documentation_block:
```


ID AAAA28525 standard; cDNA; 2348 BP.

XX
AC AAA28525;

DT 29-AUG-2000 (first entry)

DE Human oploid growth factor receptor cDNA spliced version 4.

XX
KW OGFfr: oploid growth factor receptor; growth inhibitor; proliferative;
OS cytosstatic; vulinerary. gene therapy; antagonist; chromosome 20q13.3; ss.
XX Homo sapiens.

XX
FH Key Location/Qualifiers

FT 5'UTR 1..33 /*tag= a

FT CDS 34..2067 /*tag= b

FT FT /product= Oploid_growth_factor_receptor

FT 3'UTR 2068..2348 /*tag= c

PN MO2000026340-A2.

PD 11-MAY-2000.

XX 02-NOV-1999; 99WO-US25802.

PF XX 03-NOV-1998; 98US-0106879.

PR (PENN-) PENN STATE RES FOUND.

XX
PI Zagon IS, McLaughlin PJ, Verderame MF;

XX WPI: 2000-365594/31.

DR P-P-SDB; AA92811.

PT New cDNA encoding rat and human oploid growth factor receptors which
modulate cell growth, useful for treating cancer

PS
XX
XX Claim 1; Page 84-86; 91pp; English.

CC Primers generated from rat oploid growth factor receptor (OGFr) cDNA were
used to clone a fragment of the human OGFr cDNA. The complete sequence of
human OGFr was assembled with a combination of 3' and 5' RACE. 5' RACE
consistently yielded a single species of cDNA, while the 3' RACE revealed
extensive alternative splicing. The alternate splice forms were missing
the imperfect repeats or differed in the number of imperfect repeats. The
human OGFr gene chromosomal location was determined by FISH as 20q13.3.
OGFr proteins, nucleic acid molecules, antibodies, transformed cells and
expression vector are useful for detecting expression or levels of an
OGFr in a tissue. OGFr nucleic acids can be used to inhibit growth of
cells in vitro. The antisense sequences and antibodies can be used to
control growth of cells in vitro. Cell growth can be promoted by
interfering with the OGFr ligand-receptor system, especially where a
cell subject suffers from a tissue wound. Treating cancer comprises enhancing
the function of the OGFr ligand-receptor system in cancerous cells of a
patient or administering the OGFr nucleic acid to the patient.

SO Sequence 2348 BP; 485 A; 738 G; 826 G; 299 T; 0 other;

alignment_scores:

	Quality:	41.00	Length:	9
Percent Similarity:	Ratio:	4.356	Gaps:	0
100.000	Percent Identity:	77.778		

|||||
2165 CACGAGCAGCGCTTGAGGGGAGTCCG 2139

seq_name: /SID8/gcgdelta/geneseq/geneseqn/NA2000.DAT.AAA28523
seq_documentation_block:
ID AAA28523 standard; cDNA; 2408 BP.
XX
AC AAA28523;
XX
DT 29-AUG-2000 (first entry)
XX
DE Human opiod growth factor receptor cDNA spliced version 8.
XX
KW OGFR: opiod growth factor receptor; growth inhibitor; proliferative;
KM cyostatic; vulnerary. gène therapy; antagonist; chromosome 20q13.3; ss.
XX
OS Homo sapiens.

	Key	Location/Qualifiers
FH	5'UTR	1..33
FT		/tag= a
FT	CDS	34..2127
FT		/tag= b
FT		/product= Opiod.growth_factor_receptor
FT	3'UTR	2128..2408
XX		/tag= c
PN		WO200026340-AZ.
PD		11-MAY-2000.
XX		
PF	02-NOV-1999;	99WO-US25802.
XX		
PR	03-NOV-1998;	98US-O106879.
PA	(PENN-) PENN STATE RES FOUND.	
PJ	Zagou IS, McLaughlin PJ, Verderame MF;	
DR	WPI: 2000-365594/31.	
DR	P-PsDB; AAY92809.	
XX		
XX	New cDNA encoding rat and human opiod growth factor receptors which	
PT	modulate cell growth, useful for treating cancer	
PS	Claim 1; Page 79-81; 91pp; English.	
XX		

Primers generated from rat opiod growth factor receptor (OGFR) cDNA were used to clone a fragment of the human OGFR cDNA. The complete sequence of human OGFR was assembled with a combination of 3' and 5' RACE. 5' RACE consistently yielded a single species of cDNA, while the 3' RACE revealed extensive alternative splicing. The alternate splice forms were missing the imperfect repeats or differed in the number of imperfect repeats. The human OGFR gene chromosomal location was determined by FISH as 20q13.3. expression vector are useful for detecting expression or levels of an OGFR. In a tissue. OGFR nucleic acids can be used to inhibit growth of cells in vitro. The antisense sequences and antibodies can be used to promote growth of cells in vitro. Cell growth can be promoted by interfering with the OGFR ligand-receptor system, especially where a subject suffers from a tissue wound. Treating cancer comprises enhancing the function of the OGFR ligand-receptor system in cancerous cells of a patient or administering the OGFR nucleic acid to the patient.

Sequence 2408 BP; 500 A; 762 C; 844 G; 302 T; 0 other;

```

alignment_block:
US-09-444-281-27 x AAA28525/rev ..
Align seg 1/1 to reverse of: AAA28525 from: 1 to: 2348
1 HisGLUaIaGUpProGuaIaGUpPro 9

```

```
alignment_scores:
    Quality: 41.00
    Ratio: 4.556
    Percent Similarity: 100.000
    Length: 9
    Gaps: 0
    Percent Identity: 77.778
alignment_block:
```

US-09-444-281-27 x AAA28523/rev ..

Align seg 1/1 to reverse of: AAA28523 from: 1 to: 2408

1 HisGLUaIaGluProGLUaIaGluPro 9
|||||
2225 CACGAGCAGAGCCTGAGGAGGCCG 2199

seq_name: /SID58/gcgdata/geneseq/geneseqn/NA2001.DAT:AAH17441

seq_documentation_block:

ID AAH17441 standard; cDNA; 2409 BP.

AAH17441;

26-JUN-2001 (first entry)

Human cDNA sequence SEQ ID NO:16891.

Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.

Homo sapiens.

EP1074617-A2.

07-FEB-2001.

28-JUL-2000; 2000BP-0116126.

29-JUL-1999; 99JP-0248036.

27-AUG-1999; 99JP-0300253.

11-JAN-2000; 2000JP-018776.

02-MAY-2000; 2000JP-0183767.

09-JUN-2000; 2000JP-0241899.

(HELI-) HELIX RES INST.

Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;

Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

WPI; 2001-318749/34.

Claim 8; SEQ ID 16891; 2537bp + CD ROM; English.

The present invention describes primer sets for synthesizing 5602 full-length cDNAs defined in the specification. Where a primer set comprises: (a) an oligo-dT primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprises one of the 5602 nucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides; or (b) a combination of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide which comprises a 3'-end sequence, where the oligonucleotide comprises at least 15 nucleotides and the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the specification. The primer sets can be used in antisense therapy and in gene therapy. The primers are useful for synthesizing polynucleotides, particularly full-length cDNAs. The primers are also useful for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length cDNAs easily without any specialized methods. AAH03166 to AAH13628 and AAH95893 represent human amino acid sequences; and AAH13629 to AAH13632 represent oligonucleotides, all of which are used in the exemplification of the present invention.

Sequence 2409 BP; 495 A; 757 C; 842 G; 315 T; 0 other;

alignment_scores:

Quality: 41.00 Length: 9
Ratio: 4.556 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:

US-09-444-281-27 x AAH17441/rev ..

Align seg 1/1 to reverse of: AAH17441 from: 1 to: 2409

1 HisGLUaIaGluProGLUaIaGluPro 9
|||||
2157 CACGAGCAGAGCCTGAGGAGGCCG 2131

seq_name: /SID58/gcgdata/geneseq/geneseqn/NA2001.DAT:AAH18239

seq_documentation_block:

ID AAH18239 standard; cDNA; 5796 BP.

AAH18239;

26-JUN-2001 (first entry)

Human cDNA sequence SEQ ID NO:18182.

Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.

Homo sapiens.

EP1074617-A2.

07-FEB-2001.

28-JUL-2000; 2000BP-0116126.

29-JUL-1999; 99JP-0248036.

27-AUG-1999; 99JP-0300253.

11-JAN-2000; 2000JP-018776.

02-MAY-2000; 2000JP-0183767.

09-JUN-2000; 2000JP-0241899.

(HELI-) HELIX RES INST.

Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;

Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

WPI; 2001-318749/34.

Claim 8; SEQ ID 18182; 2537bp + CD ROM; English.

The present invention describes primer sets for synthesizing 5602 full-length cDNAs defined in the specification. Where a primer set comprises: (a) an oligo-dT primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprises one of the 5602 nucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides; or (b) a combination of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide which comprises a 3'-end sequence, where the oligonucleotide comprises at least 15 nucleotides and the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the specification. The primer sets can be used in antisense therapy and in gene therapy. The primers are useful for synthesizing polynucleotides, particularly full-length cDNAs. The primers are also useful for the detection and/or diagnosis of the abnormality of the proteins encoded by

CC the full-length cDNAs. The primers allow obtaining of the full-length
CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to
CC AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632
CC represent oligonucleotides, all of which are used in the exemplification
CC of the present invention.

XX
SQ Sequence 5796 BP; 1434 A; 1616 C; 1531 G; 1215 T; 0 other;

alignment_scores:
Quality: 40.00 Length: 8
Ratio: 5.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 87.500

alignment_block:
US-09-444-281-27 x AAH18239/rev ..

Align seg 1/1 to reverse of: AAH18239 from: 1 to: 5796

1 HiscGuaLagUpProGuaLagU 8
|||||
3514 CATGAGCTGAGCTCAGCTCGAA 3491

seq_name: /SID58/gcgdata/geneseq/geneseqn/NA2001.DAT:AAH18247

seq_documentation_block:
ID AAH18247 standard; cDNA: 6048 BP.

XX
AC AAH18247;
XX
XX
DT 26-JUN-2001 (first entry)
XX
DE Human cDNA sequence SEQ ID NO:18198.
XX
XX Human; primer: detection; diagnosis; antisense therapy; gene therapy; ss.
XX
XX Homo sapiens.
XX
OS
XX
PN EPI074617-A2.
XX
PD 07-FEB-2001.
XX
PF 28-JUL-2000; 2000EP-0116126.
XX
XX
PR 29-JUL-1999; 99JP-0248036.
PR 27-AUG-1999; 99JP-0300253.
PR 11-JAN-2000; 2000JP-0118776.
PR 02-MAY-2000; 2000JP-0183767.
PR 09-JUN-2000; 2000JP-0241899.
XX
XX
PA (HELI-) HELIX RES INST.
XX
XX Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX
XX
DR WPI: 2001-318749/34.
XX
XX
PT Primer sets for synthesizing polynucleotides, particularly the 5602
PT full-length cDNAs defined in the specification, and for the detection
PT and/or diagnosis of the abnormality of the proteins encoded by the
PT full-length cDNAs -
XX
XX
PS Claim 8: SEQ ID 18198; 2537PP + CD ROM; English.
XX
XX
XX The present invention describes primer sets for synthesising 5602
XX full-length cDNAs defined in the specification. Where a primer set
XX comprises: (a) an oligo-ol primer and an oligonucleotide complementary
XX to the complementary strand of a polynucleotide which comprises one of
XX the 5602 nucleotide sequences defined in the specification, where the
XX oligonucleotide comprises at least 15 nucleotides; or (b) a combination
XX of an oligonucleotide comprising a sequence complementary to the
XX complementary strand of a polynucleotide which comprises a 5'-end

CC sequence and an oligonucleotide comprising a sequence complementary to a
CC polynucleotide which comprises a 3'-end sequence, where the
CC oligonucleotide comprises at least 15 nucleotides and the combination of
CC the 5'-end sequence/3'-end sequence is selected from those defined in
CC the specification. The primer sets can be used in antisense therapy and
CC in gene therapy. The primers are useful for synthesising polynucleotides,
CC particularly full-length cDNAs. The primers are also useful for the
CC detection and/or diagnosis of the abnormality of the proteins encoded by
CC the full-length cDNAs. The primers allow obtaining of the full-length
CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to
CC AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632
CC represent oligonucleotides, all of which are used in the exemplification
CC of the present invention.

XX
SQ Sequence 6048 BP; 1488 A; 1693 C; 1604 G; 1263 T; 0 other;

alignment_scores:
Quality: 40.00 Length: 8
Ratio: 5.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 87.500

alignment_block:
US-09-444-281-27 x AAH18247/rev ..

Align seg 1/1 to reverse of: AAH18247 from: 1 to: 6048

1 HiscGuaLagUpProGuaLagU 8
|||||
3767 CATGAGCTGAGCTCAGCTCGAA 3744

seq_name: /SID58/gcgdata/geneseq/geneseqn/NA2001.DAT:AAI61181

seq_documentation_block:
ID AAI61181 standard; cDNA: 8605 BP.

XX
AC AAI61181;
XX
XX
DT 22-OCT-2001 (first entry)
XX
XX
DE Human polynucleotide SEQ ID NO 5170.
XX
XX
XX Human; nootropic; immunosuppressant; cytostatic; gene therapy; cancer;
XX peripheral nervous system; neuropathy; central nervous system; CNS;
XX Alzheimer's; Parkinson's disease; Huntington's disease; haemostatic;
XX amyotrophic lateral sclerosis; Shy-Drager Syndrome; Chemotactic;
XX chemokinetic; thrombolytic; drug screening; arthritis; inflammation;
XX leukemia; ss.
XX
XX
XX Homo sapiens.
XX
XX
PN WO200153312-A1.
XX
XX
PD 26-JUL-2001.
XX
XX
PF 26-DEC-2000; 2000WO-US34263.
XX
XX
PR 21-JAN-2000; 2000US-0488725.
PR 25-APR-2000; 2000US-0552317.
PR 09-JUL-2000; 2000US-0598042.
PR 19-JUL-2000; 2000US-0620312.
PR 03-AUG-2000; 2000US-0653450.
PR 14-SEP-2000; 2000US-0662191.
PR 19-OCT-2000; 2000US-0693036.
PR 29-NOV-2000; 2000US-0727344.
XX
XX
XX (HYSE-) HYSEQ INC.
XX
XX
XX Tang YT, Liu C, Asundi V, Chen R, Ma Y, Qian XB, Ren F, Wang D;
PI Wang J, Wang Z, Wehrman T, Xu C, Xue AJ, Yang Y, Zhang J;
PI Zhao QA, Zhou P, Goodrich R, Drmanac RT;
XX

DR WPI: 2001-442253/47.
DR P-PSDB; AAM42025.
XX Novel nucleic acids and polypeptides, useful for treating disorders
PT such as central nervous system injuries -
XX
PS Claim 1: SEQ ID NO 5170; 10078bp; English.
XX
CC The invention relates to human nucleic acids (AA157798-AA161369) and
CC the encoded polypeptides (AAM38642-AAM42213) with neurotrophic,
CC immunosuppressant and cytoskeletal activity. The polynucleotides are useful
CC in gene therapy. A composition containing a polypeptide or polynucleotide
CC of the invention may be used to treat diseases of the peripheral nervous
CC system, such as peripheral nervous injuries, peripheral neuropathy and
CC Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic
CC lateral sclerosis, and Shy-Drager Syndrome. Other uses include the
CC utilisation of the activities such as: Immune system suppression,
CC Activin/inhibin activity, chemotactic/chemokinetic activity, haemostatic
CC and thrombolytic activity, cancer diagnosis and therapy, drug screening,
CC C.N.S disorders.
CC Note: The sequence data for this patent did not form part of the printed
CC specification.
XX
SQ Sequence 8605 BP; 2079 A; 2209 C; 2192 G; 2125 T; 0 other;
XX
alignment_scores:
Quality: 40.00 Length: 8
Ratio: 5.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 87.500
XX
alignment_block:
US-09-444-281-27 x AA161181 ..
XX
Align seg 1/1 to: AA161181 from: 1 to: 8605
XX
1 H1SGTUA1AG1UPROG1UA1AG1U 8
|||||
5995 CATGAGCTGAGCCTCAGCGTGAA 6018
XX
seq_name: /SIDS8/gcgdata/geneseq/geneseqn/NA2001.DAT:AA159395
XX
seq_documentation_block:
ID AA159395 standard; cDNA; 8840 BP.
XX
AC AA159395;
XX
DT 22-OCT-2001 (first entry)
XX
DE Human polynucleotide SEQ ID NO 1598.
XX
KW Human; neurotrophic; immunosuppressant; cytoskeletal; gene therapy; cancer;
KW peripheral nervous system; neuropathy; central nervous system; CNS;
KW Alzheimer's; Parkinson's disease; Huntington's disease; haemostatic;
KW amyotrophic lateral sclerosis; Shy-Drager Syndrome; chemotactic;
KW chemokinetic; thrombolytic; drug screening; arthritis; inflammation;
KW leukaemia; ss.
XX
OS Homo sapiens.
XX
PN WO200153312-A1.
XX
PD 26-JUL-2001.
XX
PE 26-DEC-2000; 2000WO-US34263.
XX
PR 21-JAN-2000; 2000US-0488725.
PR 25-APR-2000; 2000US-0552317.
PR 09-JUL-2000; 2000US-0588042.
PR 19-JUL-2000; 2000US-0620312.
PR 03-AUG-2000; 2000US-0653450.
XX

PR 14-SEP-2000; 2000US-06623191.
PR 19-OCT-2000; 2000US-0693036.
PR 29-NOV-2000; 2000US-0727344.
XX
XX (HYSE-) HYSEQ INC.
XX
PI Tang YT, Liu C, Asundi V, Chen R, Ma Y, Qian XB, Ren F, Wang D;
PI Wang J, Wang Z, Wehrman T, Xu C, Xue AJ, Yang Y, Zhang J;
PI Zhao QA, Zhou P, Goodrich R, Dirmacac RT;
XX
DR WPI: 2001-442253/47.
DR P-PSDB; AAM40239.
XX
PT Novel nucleic acids and polypeptides, useful for treating disorders
PT such as central nervous system injuries -
XX
PS Claim 1: SEQ ID NO 1598; 10078bp; English.
XX
CC The invention relates to human nucleic acids (AA157798-AA161369) and
CC the encoded polypeptides (AAM38642-AAM42213) with neurotrophic,
CC immunosuppressant and cytoskeletal activity. The polynucleotides are useful
CC in gene therapy. A composition containing a polypeptide or polynucleotide
CC of the invention may be used to treat diseases of the peripheral nervous
CC system, such as peripheral nervous injuries, peripheral neuropathy and
CC Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic
CC lateral sclerosis, and Shy-Drager Syndrome. Other uses include the
CC utilisation of the activities such as: Immune system suppression,
CC Activin/inhibin activity, chemotactic/chemokinetic activity, haemostatic
CC and thrombolytic activity, cancer diagnosis and therapy, drug screening,
CC C.N.S disorders.
CC Note: The sequence data for this patent did not form part of the printed
CC specification.
XX
SQ Sequence 8840 BP; 2118 A; 2331 C; 2255 G; 2136 T; 0 other;
XX
alignment_scores:
Quality: 40.00 Length: 8
Ratio: 5.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 87.500
XX
alignment_block:
US-09-444-281-27 x AA159395/rev ..
XX
Align seg 1/1 to reverse of: AA159395 from: 1 to: 8840
XX
1 H1SGTUA1AG1UPROG1UA1AG1U 8
|||||
3604 CATGAGCTGAGCCTCAGCGTGAA 3581
XX
seq_name: /SIDS8/gcgdata/geneseq/geneseqn/NA1998.DAT:AAV68396
XX
seq_documentation_block:
ID AAV68396 standard; cDNA to mRNA; 9408 BP.
XX
AC AAV68396;
XX
DT 05-MAY-1999 (first entry)
XX
DE Human BA22-alpha cDNA.
XX
KW Transcriptional regulator; transcription; BA21-alpha; bromodomain; BA2;
KW atypical zinc finger; testis; human; tumour; BA21-beta; BA22-alpha; drug;
KW BA22-beta; treatment; cancer; proliferative disorder; screening; ds.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FH CDS 740..6376
FT /*tag= a
FT /product= "BA22-alpha"

XX MO9847920-A1.
 PN
 XX
 PD 29-OCT-1998.
 XX
 PF 17-APR-1998; 98MO-JP01783.
 XX
 PR 24-OCT-1997; 97JP-0310027.
 PR 18-APR-1997; 97JP-0116570.
 XX
 PA (CHUG-) CHUGAI RES INST MOLECULAR MEDICINE INC.
 XX
 PI Jones MH;
 XX
 DR WPI: 1998-583603/49.
 DR P-PSDB; AAW81170.
 XX
 PT Transcriptional regulator gene family containing bromodomain - may
 PT be expressed in testis tissue and is useful for treatment of cancer
 PT and other proliferative disorders
 XX
 PS Claim 2; Page 72-88; 187pp; Japanese.
 XX
 CC This sequence encodes the human BAZ2-alpha protein, a member of a
 CC family of transcriptional regulator genes containing a bromodomain (BAZ,
 CC Bromodomain with Atypical zinc finger) which are expressed specifically
 CC in testis tissue and also in certain tumour lines. Transgenic cells may
 CC be used for the preparation of the BAZ1-alpha, BAZ1-beta, BAZ2-alpha and
 CC BAZ2-beta proteins. These proteins can be used in the treatment of cancer
 CC and other proliferative disorders, and in screening of compounds for
 CC their binding ability to the expression products (e.g. for use as drugs
 CC by modulation of transcriptional regulation).
 XX
 SQ Sequence 9408 BP; 2279 A; 2458 C; 2383 G; 2280 T; 8 other;

alignment_scores:
 Quality: 40.00 Length: 8
 Ratio: 5.000 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 87.500

alignment_block:
 US-09-444-281-27 x AAV68396/rev ..
 Align seg 1/1 to reverse of: AAV68396 from: 1 to: 9408
 1 HtsgtAaIaGtUpGtAaIaGtU 8
 |||||
 4343 CATGAAGCTGAGCTCAGGCTGAA 4320

THIS PAGE BLANK (USPTO)

zebrafish.
Danio rerio

REFERENCE
AUTHORS

Clark, M., Johnson, S. L., Lehrach, H., Lee, R., Li, F., Maria, M., Fady,

TITLE	JOURNAL	COMMENT
Washu Zebrafish EST Project 1998	Unpublished (1998)	Other_ESTs: f384f04.x1

Contact: Stephen L. Johnson
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: Zbratishewatson.wustl.edu
cDNA Library Preparation: Leonard Ira Zon DNA Sequencing by:
Washington University Genome Sequencing Center Clone distribution:
Genome Systems, St. Louis, Missouri (web address:
www.genomesystems.com) (email contact:
info@genomesystems.com) and
Research Genetics, Huntsville, Alabama (web address:
www.resgen.com) (email contact:
info@resgen.com) and
ResourceZentrumPrimatDatenbank, Berlin, Germany (web address:
www.izpd.de)

FEATURES	
Source	
	Seq primer: T3 ET from Amersham
	High quality sequence stop: 421.
	Location/Qualifiers
	1..593

```

/organism="Danio rerio"
/db_xref="taxon:7955"
/collection="zebrafish gridded kidney"
/sex="mixed"
/tissue_type="kidney pooled from 300 wild type adults"
/lab_host="XLOLR"
/notice="Organ: kidney; Vector: pRK-CMV; Site_1: EcoRI; Site_2: XhoI; Oligo dT cDNA library constructed from mRNA pooled from pooled kidney tissue from 300 adult zebrafish."
BASE COUNT
138 a 153 c 181 g 121 t
ORIGIN

```

```

alignment_scores:
  Quality: 45.00
  Ratio: 4.500
  Percent Similarity: 90.909
  Length: 11
  Gaps: 0
  Percent Identity: 63.636

```

```

Alignment_block:
US-09-444-281-27 x AW419871/rev
..
Align seg 1/1 to reverse of: AW419871 from: 1 to: 593

```

```

1 HisGluAlaGluProGluAlaGluProIleMet 11
  |||||:||||||| |||||:
276 CACGAGTCGAGCCAGAACCAACCTTTGGTG 244
eq_name: gb_gss:CNS01L9U

```

seq_documentation_block:	848 bp	DNA	GSS	14-JUN-2001
LOCUS	CNS01199			
DEFINITION	Anopheles gambiae GSS r7 end of clone 18M20 of NotreDame1 library from strain PST of Anopheles gambiae (African malaria mosquito), genomic survey sequence.			
ACCSSION				

RELEASED
 VERSION
 EYWORDS
 SOURCE
 ORGANISM
 ALL49363
 ALL49363.1
 GI:7009842
 GSS.
 African malaria mosquito.
 Anopheles gambiae

REFERENCE
1 (bases 1 to 848)

AUTHORS
 TITLE
 JOURNAL
 Direct Submission
 Submitted (16-FEB-2000) Genoscope - Centre National de Sequencage
 BP 101 91006 EVRY cedex - FRANCE (E-mail : sequest@genoscope.cns.fr)
 Web : www.genoscope.cns.fr
 2 (bases 1 to 848)
 REFERENCE
 PubMed

AUTHORS
 TITLE
 JOURNAL
 Submitted (16-FEB-2000) BBMI, Institut Pasteur, 25, rue du Dr

COMMENT: This clone is from an A. gambiae BAC library provided by F. H. Collins and sequenced by Genoscope in collaboration with the Laboratory of Biochem. and Biol. Molec. of Insects, Institut Pasteur.

Source	Location/Qualifiers
1.	.848

/organism="Anopheles gambiae"
 /strain="PEST"
 /db_xref="taxon:7165"
 /clone="18M20"
 /clone_1b="Notredame1"
 /note="end : T7"
 216 a 157 c 204 g 248 t 23 others

```

alignment_scores:
  Quality: 45.00
  Ratio: 4.500
  Percent Similarity: 90.909
  Length: 11
  Gaps: 0
  Percent Identity: 72.727

```

alignment_block:
US-09-444-281-27 x CNS01L9U/rev .

Align seg 1/1 to reverse of: CNS01L9U from: 1 to: 848

```

1 H1sgluAlagluProgluAlagluProIleMet 11
  |||||::: |||||:::|||||||
666 CATGATCCATACCTGATCGGACCTCATG 634

```

seq_name: gb_gss:AZ208017

```
seq_documentation_block: 0001
docline 00000017
```

Accession	A220801	GSS	31-MHG-2000
Definition	SP.0134.B1.D06.773 Strongylocentrotus purpuratus, purple sea urchin / spectrum genomic BAC library Strongylocentrotus purpuratus genomic clone Plate=134 Col=11 Row=H, DNA sequence.		
Accession	A2208017		

VERSION	AZ208017.1	GI:8420201
KEYWORDS	GSS.	
SOURCE	Strongylocentrotus purpuratus	
ORGANISM	Strongylocentrotus purpuratus	

REFERENCE
AUTHORS

Eukaryota; Metazoa; Echinodermata; Eleutherozoa; Echinozoa
Echinoidea; Euechinoidea; Echinacea; Echinoida;
Strongylocentrotidae; Strongylocentrotus.
1 (bases 1 to 923)

Cameron, R.A., Mahlreis, G., Rast, J.P., Martinez, P., Biondi, T.

Swartzell, S., Wallace, J. C., Poustka, A. J., Livingston, B. T., Wray, G. A., Ettensohn, C. A., Lehrach, H., Britten, R. J., Davidson, E. H. and Hood, L.

TITLE	assa urchin genome project: Sequence scan, virtual map, and additional resources
JOURNAL	Proc. Natl. Acad. Sci. U. S. A. 97 (17), 9514-9518 (2000)
MEDLINE	20402566
COMMENT	Contact: Cameron, RA, Davidson, EH, Hood, L

Division of Biology 156-29
California Institute of Technology
Pasadena California 91125, USA
Tel: (626) 395-8421

Fax: (626) 793-3047
Email: acamerone@caltech.edu
Plate: 134 row: 18 column: 11
Seq primer: T7
Class: BAC ends
High quality sequence stop: 923.
Location/Qualifiers

FEATURES
source

1. 923
/organism="Strongylocentrotus purpuratus"
/db_xref="taxon:7668"
/clone="plate-134 col-11 Row-H"
/clone_lib="Strongylocentrotus purpuratus, purple sea
urchin, sperm genomic BAC library"
/note="Organ: sperm; Vector: BAC3.6; BAC clones in E-Coli
DH10B"

BASE COUNT 173 a 216 c 203 g 331 t
ORIGIN

alignment_scores:

Quality: 45.00 Length: 9
Ratio: 5.625 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 88.889

alignment_block:

US-09-444-281-27 x AZ208017/rev

Align seg 1/1 to reverse of: AZ208017 from: 1 to: 923

1 HsGluAlaGluProGluAlaGluPro 9
|||||
55 CACGAAGCCGAGCCAGCCAGACCA 29

seq_name: gb_est2:BF525048

seq_documentation_block:

LOCUS BF525048 343 bp mRNA EST 11-DEC-2000
DEFINITION UI-R-AD0-vz-d-06-0-UI-r1 UI-R-AD0 Rattus norvegicus cDNA clone
UI-R-AD0-vz-d-06-0-UI 5', mRNA sequence.
ACCESSION BF525048
VERSION BF525048.1 GI:11633015
KEYWORDS EST.

SOURCE

ORGANISM Norway rat.
Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

REFERENCE 1 (bases 1 to 343)
AUTHORS Bonaldo,M.F., Lennon,G. and Soares,M.B.
TITLE Normalization and subtraction: two approaches to facilitate gene
discovery

JOURNAL Genome Res. 6 (9), 791-806 (1996)
MEDLINE 97044477
COMMENT Contact: Soares, MB.
Program for Rat Gene Discovery and Mapping
University of Iowa
451 Eckstein Medical Research Building Iowa City, IA 52242, USA
Tel: 319 335 8250
Fax: 319 335 9565
Email: msoares@blue.weeg.uiowa.edu
CDNA Library Preparation: M.B. Soares Lab Clone distribution:
Clones will be available through Research Genetics (www.resgen.com)
This clone is also available through the I.M.A.G.E. Consortium at
LNL (info@image.lnl.gov). IMAGE ID= 1794853
Seq primer: M13 Forward.

FEATURES

source

Location/Qualifiers
1. 343
/organism="Rattus norvegicus"
/strain="Sprague-Dawley"
/db_xref="taxon:10116"
/clone="UI-R-AD0-vz-d-06-0-UI"
/clone_lib="UI-R-AD0"
/dev_stage="adult"

/lab_host="DH10B (Life Technologies)"
/note="Vector: pTZ19D-Pac (Pharmacia) with a modified
polylinker. Site 1: Not I; Site 2: Eco RI; The UI-R-AD0
library is a non-normalized library constructed from 15
dpc rat atrium. The tag is a string of 5 nucleotides
present between the Not I site and the 0190-01 track.
The library was constructed as described by Bonaldo,
Lennon and Soares, Genome Research 6: 791-806, 1996.
Tissue provided by Jim Lin, Department of Biology,
University of Iowa."

BASE COUNT 106 a 83 c 95 g 59 t
ORIGIN

alignment_scores:

Quality: 43.00 Length: 11
Ratio: 3.909 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 63.636

alignment_block:

US-09-444-281-27 x BF525048

Align seg 1/1 to: BF525048 from: 1 to: 343

1 HsGluAlaGluProGluAlaGluProIleMet 11
|||||
127 CATCATGCAGACGCTCATGCAGACCTCTTGC 159

seq_name: gb_est1:AW140836

seq_documentation_block:

LOCUS AW140836 552 bp mRNA EST 30-OCT-1999
DEFINITION EST290918 Normalized rat heart, Bento Soares Rattus sp. cDNA clone
RG1AV22 5' end similar to peroxisomal farnesylated protein, mRNA
sequence.
ACCESSION AW140836
VERSION AW140836.1 GI:6160674
KEYWORDS EST.

SOURCE

ORGANISM Rattus sp.
Rattus sp.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

REFERENCE 1 (bases 1 to 552)
AUTHORS Lee,N.H., Glodok,A., Chandra,I., Mason,T.M., Quackenbush,J.,
Kerlavage,A.R. and Adams,M.D.
TITLE Rat Genome Project: Generation of a Rat EST (RESt) Catalog & Rat
Gene Index
JOURNAL Unpublished (1998)
COMMENT Other ESTs: TC85992

JOURNAL
COMMENT

The Institute for Genomic Research
9712, Medical Center Drive, Rockville, MD 20850, USA
Tel: (301)-838-3529
Fax: (301)-838-0208
Email: nhlee@tigr.org
For clone availability, additional sequence and expression
information related to this EST please check the TIGR Rat Gene
Index (http://www.tigr.org/tdb/t91/t91.html). To order a clone
contact the ATCC (http://www.atcc.org/atcc.html).
Seq primer: M13 Reverse.

FEATURES

source

Location/Qualifiers
1. 552
/organism="Rattus sp."
/db_xref="taxon:10118"
/clone="RG1AV22"
/clone_lib="Normalized rat heart, Bento Soares"
/note="Organ: heart; Vector: pTZ19D-Pac; Site 1: EcoRI;
Site 2: NotI"

BASE COUNT 155 a 138 c 149 g 110 t
ORIGIN


```

/lab.host="TJ0121"
/note="vector: lambdaZAP; Site_1: EcoRI; Site_2: XhoI. For more details on library preparation and sequence analysis see http://www.genome.clemson.edu/projects/barley/ To order a clone see http://www.genome.clemson.edu/orders"
BASE COUNT      163 a      233 c      275 g      165 t
ORIGIN

alignment_scores:
    Quality:      43.00          Length:      9
    Ratio:        4.778         Gaps:      0
Percent Similarity: 100.000     Percent Identity: 77.778

alignment_block:
US-09-444-281-27 x BE259969 ..

Align seg 1/1   to: BE259969 from: 1   to: 836

1 HlSGluAlagluProGluAlagluPro 9
|||||:|||||||:|||||||
614 CATGATCTGTGAACCGAAGCTAGCCG 640

seq_name: gb_est1.BE037128

seq_documentation_block:
LOCUS       BE037128      1247 bp      mRNA           EST       07-JUN-2000
DEFINITION  Mp1SE02 MP Mesembryanthemum crystallinum cDNA 5' similar to
            auxin-regulated protein, mRNA sequence.
ACCESSION   BE037128
VERSION     BE037128.1 GI:8332144
KEYWORDS    EST.
SOURCE      common ice plant.
ORGANISM    Mesembryanthemum crystallinum
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
            Caryophyllidae; Caryophyllales; Alzaceae; Mesembryanthemum.
REFERENCE   1 (bases 1 to 1247)
            Bonnet,H.J., Borchert,C., Brazille,S., Brooks,J., Eaton,M., Ferreira
            ,H., Kawasaki,S., McCollough,A., Michalowski,C.B., Palacios,C.,
            Scara,G., Wheeler,M. and Zepeda,G.R.
            Functional Genomics of Plant Stress Tolerance
            Unpublished (2000)
COMMENT     Contact: Michalowski,C.B.
            University of Arizona
            Bio Sciences West room 513, Tucson, AZ 85721, USA
            Tel: 520-621-7982
            Fax: 520-621-1697
            Email: cbm@u.arizona.edu.
FEATURES             Location/Qualifiers
     source          1..1247
                     /organism="Mesembryanthemum crystallinum"
                     /db_xref="taxon:3544"
                     /clone_lib="MP"
                     /tissue_type="apical meristem and leaf primordia"
                     /dev_stage="6 weeks"
                     /note="3 d 500mm NaCl"
BASE COUNT      270 a      300 c      242 g      301 t      134 others
ORIGIN

alignment_scores:
    Quality:      43.00          Length:      10
    Ratio:        5.375         Gaps:      0
Percent Similarity: 80.000     Percent Identity: 70.000

alignment_block:
US-09-444-281-27 x BE037128/rev ..

Align seg 1/1   to reverse of: BE037128 from: 1   to: 1247

1 HlSGluAlagluProGluAlagluPro10
||||| ||||||| |||||:|:|

```

639 CATGACCCGACCGACCGACCGACCGTC 610

seq_name: gb_est2:BF655720

seq_documentation_block:

LOCUS BF655720 181 bp mRNA EST 20-DEC-2000
DEFINITION Fm1_48_B05_b1_A003 Floral-Induced Meristem 1 (Fm1) Sorghum
PROPINQUUM cdna, mRNA sequence.
ACCESSION BF655720
VERSION BF655720.1 GI:11920852
KEYWORDS EST

SOURCE Sorghum propinquum.

ORGANISM Sorghum propinquum.
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC
clade; Panicoideae; Andropogoneae; Sorghum.
1 (bases 1 to 181)
Cordonnier-Pratt,M.-M., Gingle,A., Sudman,M., Marsala,C. and Pratt,
L.H.

REFERENCE An EST database from Sorghum: floral-induced meristems
AUTHORS Unpublished (2000)
COMMENT Contact: Cordonnier-Pratt MM
Department of Botany
The University of Georgia
Plant Sciences Building, Rm. 2502, Athens, GA 30602-7271, USA
Tel: 706 542 1860
Fax: 706 542 1805
Email: mpratt@ga.edu

Sequences have been trimmed to exclude Polya, vector and regions
below Phred quality 16. The threshold for highest quality sequence
is 20.

Seq primer: JEN REV
High quality sequence stop: 111
POLYA-NO.

FEATURES
Source Location/Qualifiers

1..181
/organism="Sorghum propinquum"
/db_xref="taxon:132711"
/clone_lib="Floral-Induced Meristem 1 (Fm1)"
/note="Origin: Floral-Induced Meristems; Vector:
pJuescript II; From Lambda Zap II; Site 1: XhoI; Site 2:
EcoRI; mature plants were placed in a growth chamber for
15 days with 16 hr darkness and 8 hr light (flowering is
induced by short-day conditions); 16 days after being
returned to the greenhouse under natural long days during
late April/early May, meristems were harvested. The
library was made from poly-A RNA in the cloning vector
lambda Zap II. Clones to be sequenced were prepared by
mass excision."
BASE COUNT 52 a 40 c 61 g 28 t
ORIGIN

alignment_scores:
Quality: 42.00 Length: 11
Ratio: 4.200 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:
US-09-444-281-27 x BF655720

Align seg 1/1 to: BF655720 from: 1 to: 181

1 HisGUAAGAGTuproGUAAGAGTuproLleMet 11
|||||:|||||:|||||:|||||:|||||:|||||:
107 CATCAGCGCGACCGACCGACCGACCGATCCTA 139

seq_name: gb_est2:T70690

seq_documentation_block:

LOCUS T70690 323 bp mRNA EST 17-OCT-1996
DEFINITION 84 vegetative meristem Zea mays cdna clone 7C03B11, mRNA sequence.
ACCESSION T70690

VERSION T70690.1 GI:681838
KEYWORDS EST.
SOURCE Zea mays.
ORGANISM Zea mays

REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC
clade; Panicoideae; Andropogoneae; Zea.
1 (bases 1 to 323)
Shen,B., Carneiro,N., Torres-Jerez,I., Stevenson,R., McCreery,T.,
Helentjaris,T., Baysdorfer,C., Almira,E., Ferl,R., Habbema,D. and
Larkin,B.

TITLE Partial sequencing and mapping of clones from two maize CDNA
JOURNAL libraries
MEDLINE Plant Mol. Biol. 26, 1085-1101 (1994)
COMMENT 95111093
Contact: The Maize CDNA Project

Helentjaris TG (primary contact)
Pioneer Hi-Bred Int., Inc.
Agronomic Traits/TTD
7300 N.W. 62nd Ave. P.O. Box 1004
Johnston, IA 50131-1004
ph: 515-270-3691
fax: 515-270-4312
E-mail: helnjartg@phbred.com

Chris Baysdorfer
Department of Biological Sciences, School of Science
California State University, Hayward
Hayward, CA 94542
ph: 510-881-3459
fax: 510-727-2035
E-mail: cbaysdor@s1.csuhayward.edu

Rob Ferl
Interdisciplinary Center for Biotechnology Research
DNA Sequencing Core
University of Florida
P.O. Box 100695
Gainesville, FL 32611-0695
ph: 904-392-1928, ext. 301
fax: 904-392-4072
E-mail: robferl@nervm.nerdc.ufl.edu

FEATURES
Source Location/Qualifiers

1..323
/organism="Zea mays"
/strain="W64A2"
/db_xref="taxon:4577"
/clone_lib="7C03B11"
/clone_lib="vegetative meristem"
/lab_host="SOLR"
/note="Vector: Zap; Site 1: EcoRI; Site 2: XhoI; ds-cDNA
was prepared from oligo-dT selected mRNA by priming with a
XhoI oligo-dT oligomer and then adding the second strand
to RNase-nicked DNA:RNA hybrid with DNA POLI. EcoRI
adaptors were added to the ends, the ds-cDNAs were then
digested with EcoRI- XhoI and size-selected. These were
directionally-cloned into the Zap phage vector, excised as
plasmids, and then individually analyzed."
BASE COUNT 55 a 91 c 97 g 64 t 16 others
ORIGIN

alignment_scores:
Quality: 42.00 Length: 10
Ratio: 5.250 Gaps: 0
Percent Similarity: 80.000 Percent Identity: 70.000

alignment_block:
US-09-444-281-27 x T70690/rev

Align seg 1/1 to reverse of: T70690 from: 1 to: 323

1 H18GUALAGUPROGLUAGLUPRO 10
||||| ||||||| |||||||
36 CACGACCAACCACTGACGACGACGCTG 7

seq_name: gb_est1:BI240337

seq_documentation_block: 393 bp mRNA EST 12-JUL-2001

LOCUS BI240337

DEFINITION R337162.5prime RE Drosophila melanogaster normalized Embryo p1c-1

Phan0007611 located on: 3L 78B1-78B1; 05/12/2001, mRNA sequence.

ACCESSION BI240337

VERSION BI240337.1 GI:14708940

KEYWORDS EST

SOURCE fruit fly.

ORGANISM Drosophila melanogaster

REFERENCE Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;

Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;

Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

1 (bases 1 to 393)

Scapleton,M., Brokstein,P., Hong,L., Tyler,D., Berman,B., Carlson

,J., Champe,M., Chavez,C., Dorsett,V., Farfan,D., Frise,E., George

,R., Gonzalez,M., Guarin,H., Harris,N., Li,P., Liao,G., Misra,S.,

Mungall,C.J., Nunoo,J., Pacleb,J., Paragas,V., Park,S.,

Phouenavong,S., Wan,K., Yu,C., Lewis,S.E., Celniker,S. and Rubin

,G.M.

BDGP/HMMI RE Drosophila EST Project

Unpublished (2001)

Contact: Stapleton, M.

BDGP

Lawrence Berkeley National Lab

One Cyclotron Rd, Berkeley, CA 94720, USA

Fax: 510 486 6798

Email: http://www.fruitfly.org/EST, est@fruitfly.berkeley.edu

hit genomic AE003594; arm:3L (21112610,21418125]

estimated-cyto:78C7-78E3; 05/12/2001

plate: RE 371 row: F column: 2

High quality sequence stop: 342.

Location/Qualifiers

1..393

/organism="Drosophila melanogaster"

/db_xref="taxon:7227"

seq_documentation_block:

LOCUS AU052906 419 bp mRNA EST

DEFINITION AU052906 Dictyostelium discoideum SL (H.Urushihara) Dictyostelium

discoideum cDNA clone SLF303, mRNA sequence.

ACCESSION AU052906

VERSION AU052906.1 GI:4701389

KEYWORDS EST

SOURCE Dictyostelium discoideum.

ORGANISM Dictyostelium discoideum.

REFERENCE Eukaryota; Mycetozoa; Dictyostelida; Dictyostelium.

1 (bases 1 to 419)

Morio,T., Urushihara,H., Saito,T., Ugawa,Y., Mizuno,H., Yoshida,M.,

Yoshino,R., Mieda,B.N., Li,M., Saito,T., Takemoto,K., Yasukawa,H.,

Williams,J., Mehta,M., Takeuchi,I., Ochiai,H. and Tanaka,Y.

Developmental cDNA in Dictyostelium discoideum

Unpublished (1998)

Contact: Hideko Urushihara

Institute of Biological Sciences

University of Tsukuba

3-3-10 Ten-noda, Tsukuba, Ibaraki 305, Japan

Email: d402huesakura.cc.tsukuba.ac.jp

PROJECT = Dictyostelium discoideum cDNA project in Japan.

Location/Qualifiers

1..419

/organism="Dictyostelium discoideum"

/strain="AX4"

/db_xref="taxon:44689"

/clone="SLF303"

/clone_lib="Dictyostelium discoideum SL (H.Urushihara)"

/dev_stage="slug"

BASE COUNT 118 a 86 c 89 g 126 t

ORIGIN

alignment_scores:

Quality: 42.00 Length: 8

Ratio: 5.250 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-27 x AU052906/rev

Align seg 1/1 to reverse of: AU052906 from: 1 to: 419

2 GUUAGLUPROGLUAGLUPRO 9

||||| ||||||| |||||||

164 GAGGCGAACCTGACGACGACCT 141

seq_name: gb_est1:AU923872

seq_documentation_block:

LOCUS AU923872 435 bp mRNA EST

DEFINITION WS1_30_B09.D1_A002 Water-stressed 1 (WS1) Sorghum bicolor cDNA,

mRNA sequence.

ACCESSION AU923872

VERSION AU923872.1 GI:8089697

KEYWORDS EST

SOURCE sorghum.

ORGANISM sorghum bicolor

REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PAC

clade; Panicoidae; Andropogoneae; Sorghum.

1 (bases 1 to 435)

Cordonnier-Pratt,M.-M., Gingle,A., Marsala,C., Sudman,M. and Pratt

,L.H.

An EST database from Sorghum: water-stressed plants

Unpublished (2000)

Contact: Cordonnier-Pratt MM

Department of Botany

The University of Georgia

Plant Sciences Building, Rm. 2502, Athens, GA 30602-7271, USA

Tel: 706 542 1860

2 GUUAGLUPROGLUAGLUPRO 9

||||| ||||||| |||||||

190 GAGGCTGAGCCTGAGGACGACCT 213

seq_name: gb_est1:AU052906

Fax: 706 542 1805
Email: mmprratt@uga.edu
Sequences have been trimmed to exclude POLYA, vector and regions
below Phred quality 16. The threshold for highest quality sequence
is 20.

Seq primer: JEN REV
High quality sequence stop: 368
POLYA-No.

FEATURES
source

1.435 location/Qualifiers
/organism="Sorghum bicolor"
/db_xref="taxon:4558"
/clone_lib="Water-stressed 1 (WS1)"
/note="Organ: Mix of 5-week old plants on days 7 & 8 after
water was withheld; Vector: Lambda Zap; Site_1: XhoI;
Site_2: EcoRI; The library was made from poly-A RNA in the
cloning vector Lambda Zap II. Clones to be sequenced were
prepared by mass excision."
156 a 162 g 84 g 33 t
BASE COUNT
ORIGIN

alignment_scores:
Quality: 42.00 Length: 10
Ratio: 4.667 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:
US-09-444-281-27 x AW923872

Align seg 1/1 to: AW923872 from: 1 to: 435

1 HisqluAlagluProgluAlagluProile 10
|||||
2 CACGAGGCTGAGCCTCAACCTAAACCATTA 31

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:40:32 ; Search time 50.17 Seconds
(Without alignments)
32.071 Million cell updates/sec

Title: US-09-444-281-27
Perfect score: 59
Sequence: 1 HEAPEPEPIM 11

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 473505 seqs, 146272329 residues
Total number of hits satisfying chosen parameters: 473505

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: SPTEMBL_17:*
2: sp_archaea:*
3: sp_bacteria:*
4: sp_fungi:*
5: sp_human:*
6: sp_invertebrate:*
7: sp_mammal:*
8: sp_mhc:*
9: sp_organelle:*
10: sp_plant:*
11: sp_rodent:*
12: sp_virus:*
13: sp_vertebrate:*
14: sp_unclassified:*

Pred. No. is the number of results predicted, by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	42	71.2	880	5	017338 caenorhabd
2	42	71.2	885	5	09TXR9 caenorhabd
3	42	71.2	930	5	017339 caenorhabd
4	41	69.5	874	5	09VO10 caenorhabd
5	40	67.8	248	12	065200 drosophila
6	40	67.8	1380	5	09V108 african swi
7	39	66.1	251	3	09US24 schizosacch
8	39	66.1	304	5	09NAB8 caenorhabd
9	39	66.1	340	5	09NAB2 caenorhabd
10	39	66.1	408	5	096979 parophrys
11	39	66.1	417	5	09W4Y3 drosophila
12	39	66.1	435	5	09NEF8 drosophila
13	39	66.1	848	5	021489 caenorhabd
14	39	66.1	1192	5	09W475 drosophila
15	39	66.1	1277	12	008547 cercopithec
16	39	66.1	1279	12	066031 cercopithec
17	39	66.1	1458	3	09HE72 neotropora
18	39	66.1	1593	5	020207 caenorhabd
19	39	66.1	6797	2	09X993 streptomyce

20	38	64.4	150	1	09YFA9 aeropyrum p
21	38	64.4	200	4	09U0B8 homo sapien
22	38	64.4	288	3	09P3H2 neotropora
23	38	64.4	592	3	059900 cyptococcu
24	38	64.4	682	2	09A808 caulobacter
25	38	64.4	1083	2	086637 streptomyce
26	37	62.7	20	4	09UD25 homo sapien
27	37	62.7	101	6	095284 sus scrofa
28	37	62.7	125	2	09W24 streptomyce
29	37	62.7	175	10	09SYE4 arabidopsis
30	37	62.7	208	4	060937 homo sapien
31	37	62.7	248	11	063224 raltus norv
32	37	62.7	251	6	09GLN2 bos taurus
33	37	62.7	265	37	09JZP0 neisseria m
34	37	62.7	265	2	09JUS3 neisseria m
35	37	62.7	338	10	09M4K6 anthoceros
36	37	62.7	338	10	09M4K5 anthoceros
37	37	62.7	356	10	064647 arabidopsis
38	37	62.7	387	5	09VZC7 drosophila
39	37	62.7	401	5	09VZC6 drosophila
40	37	62.7	407	5	09NCB6 drosophila
41	37	62.7	439	11	09UMF5 mus musculu
42	37	62.7	439	11	09DC79 mus musculu
43	37	62.7	583	4	09UKV2 homo sapien
44	37	62.7	612	5	09U003 caenorhabd
45	37	62.7	614	4	09UKV1 homo sapien

ALIGNMENTS

RESULT	1	PRELIMINARY;	PRT;	880 AA.
ID	017338			
AC	017338			
DT	01-JAN-1998 (TEMBLrel. 05, Created)			
DT	01-JAN-1998 (TEMBLrel. 05, Last sequence update)			
DT	01-NOV-1998 (TEMBLrel. 08, Last annotation update)			
DE	T23E7.2B PROTEIN.			
GN	T23E7.2B.			
OS	Caenorhabditis elegans.			
OC	Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditioidea;			
OC	Rhabditidae; Peloderinae; Caenorhabditis.			
OX	NCBI_TaxID=6239;			
RN	[1]			
RC	SEQUENCE FROM N.A.			
RC	STRAIN-BRISTOL N2;			
RX	MEDLINE-94150718; PubMed-7906398;			
RA	Wilson R., Ainscough R., Anderson K., Baynes C., Berks M.,			
RA	Bonfield J., Burton J., Connell M., Copesey T., Cooper J., Coulson A.,			
RA	Craxton M., Dear S., Du Z., Durbin R., Favello A., Fulton L.,			
RA	Gardner A., Green P., Hawkins T., Hillier L., Jier M., Johnston L.,			
RA	Jones M., Kershaw J., Kirsten J., Laister N., Latreille P.,			
RA	Lighning J., Lloyd C., McMurray A., Mortimore B., O'Callaghan M.,			
RA	Parsons J., Percy C., Rifken L., Roopra A., Saunders D., Showkeen R.,			
RA	Smaison N., Smith A., Sonhammer E., Straden R., Sulston J.,			
RA	Thierry-Mieg J., Thomas K., Vaudin M., Vaughan K., Waterston R.,			
RA	Watson A., Weinstock L., Wilkinson-Sprout J., Wohlman P.,			
RT	"2.2 Mb of contiguous nucleotide sequence from Chromosome III of C.			
RT	elegans.";			
RL	Nature 368:32-38(1994).			
RN	[2]			
RC	SEQUENCE FROM N.A.			
RC	STRAIN-BRISTOL N2;			
RA	Latreille P., Steillies L., Elliott G.,			
RL	Submitted (OCT-1997) to the EMBL/GenBank/DBJ databases.			
RN	[3]			
RC	SEQUENCE FROM N.A.			
RC	STRAIN-BRISTOL N2;			
RA	Waterston R.,			
RL	Submitted (SEP-1997) to the EMBL/GenBank/DBJ databases.			
DR	EMBL; AF026205; AAB71258.1;			
SO	SEQUENCE 880 AA; 95398 MW; 97A8A101E8FBA1C1 CRC64;			

Page 2

RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
 RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Idegwam C.,
 RA Jallali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
 RA Kimmel B.E., Kodira C.D., Krafc C., Kravitz S., Kuip D., Lai Z.,
 RA Lasko P., Lai T., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
 RA Liu X., Matel B., McIntosh T.C., McLeod M.P., McPherson D.,
 RA Merklov G., Mishina N.V., Mobaraj C., Morris J., Moshrefi A.,
 RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
 RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacleb J.M.,
 RA Palazzo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
 RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
 RA Slater E., Spradling A.C., Stapleton M., Strong R., Sun E.,
 RA Sytkas R., Teclor C., Turner R., Venter E., Wang A.H., Wang X.,
 RA Wang Z.-Y., Wasserman D.A., Weinstein G.M., Weisenbach J.,
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
 RA Ye J., Yen R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
 RA "The genome sequence of *Drosophila melanogaster*.";
 RL Science 287:2185-2195(2000).
 DR EMBL: AEO03581: AAF51156.1:-
 DR FlyBase: FBgn0031496; CG17258.
 SQ SEQUENCE 874 AA: 103694 MW: 5F56DJCE7A01D9A CRC64:

Query Match 69.5%: Score 41; DB 5; Length 874;
 Best Local Similarity 70.0%: Pred. NO. 49;
 Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Oy 2 EAEPEAPIM 11
 Db 376 EAEPEAPIM 385

RESULT 5
 ID 065200 PRELIMINARY: PRT: 248 AA.
 AC 065200:
 DT 01-NOV-1996 (TREMblrel. 01, Created)
 DT 01-NOV-1996 (TREMblrel. 01, Last sequence update)
 DT 01-NOV-1998 (TREMblrel. 08, Last annotation update)
 DE PE248R.
 GN E248R.
 OS African swine fever virus (ASFV).
 OC Viruses; dsDNA viruses, no RNA stage; Asfarviridae;
 OC African swine fever-like viruses.
 OX NCBI_TaxID=10497;
 RN 11
 RP SEQUENCE FROM N.A.
 RC STRAIN=BA71V;
 RX MEDLINE=96036500; PubMed=7483270;
 RA Yanez R.J., Rodriguez J.F., Vinuela E.,
 RA Rodriguez J.F., Vinuela E.,
 RT "Immune protection conferred by the baculovirus-related glycoprotein
 RT of Hogogo virus (Orthomyxoviridae).";
 RL Virology 208:249-278(1995).
 RN 12
 RP SEQUENCE FROM N.A.
 RC STRAIN=BA71V;
 RX MEDLINE=94233765; PubMed=8178480;
 RA La Vega I., Gonzalez A., Blasco R., Calvo V., Vinuela E.;
 RT "Nucleotide sequence and variability of the inverted terminal
 RT repetitions of African swine fever virus DNA.";
 RL Virology 201:152-156(1994).
 RN 13
 RP SEQUENCE FROM N.A.
 RC STRAIN=BA71V;
 RX MEDLINE=90219205; PubMed=2325203;
 RA Gonzalez A., Calvo V., Almazan F., Almendral J.M., Ramirez J.C.,
 RA La Vega I., Blasco R., Vinuela E.;
 RP "Multigene families in African swine fever virus: family 360.";
 RL J. Virol. 64:2073-2081(1990).

RN 14
 RP SEQUENCE FROM N.A.
 RC STRAIN=BA71V;
 RX MEDLINE=90219204; PubMed=2325202;
 RA Almendral J.M., Almazan F., Blasco R., Vinuela E.;
 RT "Multigene families in African swine fever virus: family 110.";
 RL J. Virol. 64:2064-2072(1990).
 RN 15
 RP SEQUENCE FROM N.A.
 RC STRAIN=BA71V;
 RX MEDLINE=91134988; PubMed=1994575;
 RA Camacho A., Vinuela E.;
 RT "Protein p22 of African swine fever virus: an early structural protein
 RT that is incorporated into the membrane of infected cells.";
 RL Virology 181:251-257(1991).
 RN 16
 RP SEQUENCE FROM N.A.
 RC STRAIN=BA71V;
 RA Almazan F., Murguia J.R., Rodriguez J.M., La Vega I., Vinuela E.;
 RL J. Gen. Virol. 0:0-0(0).
 RN 17
 RP SEQUENCE FROM N.A.
 RC STRAIN=BA71V;
 RX MEDLINE=94187118; PubMed=8139051;
 RA Rodriguez J.M., Yanez R.J., Pan R., Rodriguez J.F., Salas M.L.,
 RA Vinuela E.;
 RT "Multigene families in African swine fever virus: family 505.";
 RL J. Virol. 68:2746-2751(1994).
 RN 18
 RP SEQUENCE FROM N.A.
 RC STRAIN=BA71V;
 RX MEDLINE=9346971; PubMed=8393914;
 RA Yanez R.J., Rodriguez J.M., Rodriguez J.F., Salas M.L., Vinuela E.;
 RT "African swine fever virus thymidylate kinase gene: sequence and
 RT transcriptional mapping.";
 RL J. Gen. Virol. 74:1633-1638(1993).
 RN 19
 RP SEQUENCE FROM N.A.
 RC STRAIN=BA71V;
 RX MEDLINE=94065656; PubMed=8245848;
 RA Alcant A., Angulo A., Vinuela E.;
 RT "Mapping and sequence of the gene encoding the African swine fever
 RT virion protein of M(r) 11500.";
 RL J. Gen. Virol. 74:2317-2324(1993).
 RN 110
 RP SEQUENCE FROM N.A.
 RC STRAIN=BA71V;
 RX MEDLINE=93277388; PubMed=8503790;
 RA Munoz M., Freije J.M., Salas M.L., Vinuela E., Lopez-Otin C.;
 RT "Structure and expression in E. coli of the gene coding for protein
 RT p10 of African swine fever virus.";
 RL Arch. Virol. 130:93-107(1993).
 RN 111
 RP SEQUENCE FROM N.A.
 RC STRAIN=BA71V;
 RX MEDLINE=90357780; PubMed=2389555;
 RA Blasco R., Lopez-Otin C., Munoz M., Bockamp E.O., Simon-Mateo C.,
 RA Vinuela E.;
 RT "Sequence and evolutionary relationships of African swine fever virus
 RT thymidine kinase.";
 RL Virology 178:301-304(1990).
 RN 112
 RP SEQUENCE FROM N.A.
 RC STRAIN=BA71V;
 RX MEDLINE=93281390; PubMed=8506138;
 RA Yanez R.J., Boursnell M., Nogal M.L., Yuste L., Vinuela E.;
 RT "African swine fever virus encodes two genes which share significant
 RT homology with the two largest subunits of DNA-dependent RNA
 RT polymerases.";
 RL Nucleic Acids Res. 21:2423-2427(1993).
 RN 113
 RP SEQUENCE FROM N.A.
 RC STRAIN=BA71V;

RX MEDLINE-93353606; PubMed-8102411;
 RA Rodriguez J.M., Yanez R.J., Almazan F., Vinuela E., Rodriguez J.F.;
 RT "African swine fever virus encodes a CD2 homolog responsible for the
 RL adhesion of erythrocytes to infected cells."; J. Virol. 67:5312-5320(1993).
 RN [14]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-94085774; PubMed-8262374;
 RA Yanez R.J., Rodriguez J.M., Bournell M., Rodriguez J.F., Vinuela E.;
 RT "Two putative African swine fever virus helicases similar to yeast
 RL and DFR."; Gene 134:161-174(1993).
 RN [15]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-90223993; PubMed-2327074;
 RA Lopez-Otin C., Freije J.M., Parra F., Mendez E., Vinuela E.;
 RT "Mapping and sequence of the gene coding for protein p72, the major
 RL capsid protein of African swine fever virus."; Virology 175:477-484(1990).
 RN [16]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-94123986; PubMed-8293992;
 RA Rodriguez J.M., Yanez R.J., Rodriguez J.F., Vinuela E., Salas M.L.;
 RT "The DNA polymerase-encoding gene of African swine fever virus:
 RL sequence and transcriptional analysis."; Gene 136:103-110(1993).
 RN [17]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-93327788; PubMed-8335009;
 RA Simon-Mateo C., Andres G., Vinuela E.;
 RT "Polyprotein processing in African swine fever virus: a novel gene
 RL expression strategy for a DNA virus."; EMBO J. 12:2977-2987(1993).
 RN [18]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-93233210; PubMed-8474154;
 RA Prados F.J., Vinuela E., Alcamí A.;
 RT "Sequence and characterization of the major early phosphoprotein p32
 RL of African swine fever virus."; J. Virol. 67:2475-2485(1993).
 RN [19]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-92260660; PubMed-1583732;
 RA Alcamí A., Angulo A., Lopez-Otin C., Munoz M., Freije J.M.,
 RT "Amino acid sequence and structural properties of protein p12, an
 RL African swine fever virus attachment protein."; J. Virol. 66:3860-3868(1992).
 RN [20]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-93174976; PubMed-8438592;
 RA Yanez R.J., Vinuela E.;
 RT "African swine fever virus encodes a DNA ligase."; Virology 193:531-536(1993).
 RN [21]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-93174941; PubMed-8382399;
 RA Pena L., Yanez R.J., Revilla Y., Vinuela E., Salas M.L.;
 RT "African swine fever virus guanylyltransferase."; Virology 193:319-328(1993).
 RN [22]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-95159428; PubMed-7856088;

RA Simon-Mateo C., Freije J.M., Andres G., Lopez-Otin C., Vinuela E.;
 RT "Mapping and sequence of the gene encoding protein p17, a major
 RL African swine fever virus structural protein."; Virology 206:1140-1144(1995).
 RN [23]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-92263807; PubMed-1316688;
 RA Garcia-Beato R., Freije J.M., Lopez-Otin C., Blasco R., Vinuela E.;
 RT "A gene homologous to topoisomerase II in African swine fever virus."; Virology 188:938-947(1992).
 RN [24]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-94091056; PubMed-8266720;
 RA Freije J.M., Lain S., Vinuela E., Lopez-Otin C.;
 RT "Nucleotide sequence of a nucleoside triphosphate phosphohydrolase
 Query Match 67.8%; Score 40; DB 12; Length 248;
 Best Local Similarity 63.6%; Pred. No. 20;
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
 QY 1 HEAPEPEPIM 11
 Db 227 HEEEEAPEPLI 237
 ID 09VIU8 PRELIMINARY; PRT; 1380 AA.
 AC 09VIU8;
 DT 01-MAY-2000 (TREMBLrel. 13, Created)
 DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
 DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
 DE CG10132 PROTEIN.
 GN CG10132.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
 OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 OX NCBI_Taxid=7227;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BERKELEY;
 RX MEDLINE-20196006; PubMed-10731132;
 RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
 RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,
 RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
 RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
 RA Brandon R.C., Rogers Y.-H.C., Blazer R.G., Champe M., Pfeiffer B.D.,
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,
 RA Abril J.F., Abghari A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,
 RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
 RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
 RA Borkova D., Botchan M.R., Bouck B., Brokstein P., Brotler P.,
 RA Burtis K.C., Busam D.A., Butler H., Cadieu L., Center A., Chandra I.,
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
 RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
 RA Durbin K.J., Evangelista C.C., Ferraz C., Ferrelle S., Fleischmann W.,
 RA Foster C., Gabriellian A.E., Garg N.S., Gelbart W.M., Glasser K.,
 RA Glodek A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
 RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Idegami C.,
 RA Jajani M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
 RA Lasko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
 RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
 RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
 RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacle J.M.,
 RA Palazolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,

RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
 RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
 RA Strydom R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
 Wang Z.-Y., Wasserman D.A., Weinstein G.M., Weissbach J.,
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
 RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
 RT "The genome sequence of *Drosophila melanogaster*.";
 RL Science 287:2185-2195(2000).
 DR EMBL: AE003663; AAF53816.1; -
 DR FlyBase: FBgn0032798; CG10132.
 DR InterPro: IPR001680; WD40.
 DR SMART: SM00320; WD40; 2.
 DR PROSITE: PS00678; WD_REPEATS_1; 1.
 DR PROSITE: PS50294; WD_REPEATS_REGION; 1.
 KW Repeat; WD repeat.
 SQ SEQUENCE 1380 AA; 154423 MW; C1928D066450A15B CRC64;

Query Match 67.88; Score 40; DB 5; Length 1380;
 Best Local Similarity 66.78; Pred. No. 1.2e+02;
 Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 1 AEPEAP 9
 Db 179 HDSEPTEP 187

RESULT 7
 ID Q9USZ4 PRELIMINARY; PRT; 251 AA.
 AC Q9USZ4;
 DT 01-MAY-2000 (TREMblrel. 13, Created)
 DT 01-MAY-2000 (TREMblrel. 13, Last sequence update)
 DE HYPOTHETICAL 28.0 KDA PROTEIN.
 GN SPBCL1G11.05.
 OS Schizosaccharomyces pombe (Fission yeast).
 OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
 OC Schizosaccharomycetales; Schizosaccharomycetaceae;
 OC Schizosaccharomyces.
 OX NCBI_TaxID=4896;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=9728-;
 RA Saunders D., McDougall R.C., Rajandream M.A., Barrell B.G.;
 RL Submitted (OCT-1999) to the EMBL/GenBank/DBJ databases.
 DR EMBL: ALI32717; CAB59807.1; -
 KW Hypothetical protein.
 SQ SEQUENCE 251 AA; 28010 MW; 8018F8325AC65B99 CRC64;

Query Match 66.18; Score 39; DB 3; Length 251;
 Best Local Similarity 87.58; Pred. No. 30;
 Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 2 AEPEAP 9
 Db 185 ESEPEAP 192

RESULT 8
 ID Q9NAB8 PRELIMINARY; PRT; 304 AA.
 AC Q9NAB8;
 DT 01-OCT-2000 (TREMblrel. 15, Created)
 DT 01-OCT-2000 (TREMblrel. 15, Last sequence update)
 DE Y33FAB.16 PROTEIN.
 GN Y33FAB.16;
 OS *Caenorhabditis elegans*.

OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditioidea;
 OC Rhabditidae; Peloderinae; *Caenorhabditis*.
 OX NCBI_TaxID=6239;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Smye R.;
 RL Submitted (OCT-1999) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=9069613; PubMed=9851916;
 RA none;
 RT "Genome sequence of the nematode *C. elegans*: A platform for
 RT investigating biology.";
 RL Science 282:2012-2018(1998).
 DR EMBL: ALI32949; CAB61066.1; -
 SQ SEQUENCE 304 AA; 34146 MW; 20AB91D8BB137A76 CRC64;

Query Match 66.18; Score 39; DB 5; Length 304;
 Best Local Similarity 87.58; Pred. No. 37;
 Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 3 AEPEAP 10
 Db 80 AEPEAP 87

RESULT 9
 ID Q9NAA2 PRELIMINARY; PRT; 340 AA.
 AC Q9NAA2;
 DT 01-OCT-2000 (TREMblrel. 15, Created)
 DT 01-OCT-2000 (TREMblrel. 15, Last sequence update)
 DE Y33FAB.17 PROTEIN.
 GN Y33FAB.17;
 OS *Caenorhabditis elegans*.
 OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditioidea;
 OC Rhabditidae; Peloderinae; *Caenorhabditis*.
 OX NCBI_TaxID=6239;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Smye R.;
 RL Submitted (OCT-1999) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=9069613; PubMed=9851916;
 RA none;
 RT "Genome sequence of the nematode *C. elegans*: A platform for
 RT investigating biology.";
 RL Science 282:2012-2018(1998).
 DR EMBL: ALI32949; CAB70107.1; -
 SQ SEQUENCE 340 AA; 36879 MW; 2EC01E30582C9E3A CRC64;

Query Match 66.18; Score 39; DB 5; Length 340;
 Best Local Similarity 87.58; Pred. No. 42;
 Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 3 AEPEAP 10
 Db 247 AEPEAP 254

RESULT 10
 ID O96979 PRELIMINARY; PRT; 408 AA.
 AC O96979;
 DT 01-MAY-1999 (TREMblrel. 10, Created)
 DT 01-MAY-1999 (TREMblrel. 10, Last sequence update)
 DE TRANSLATION ELONGATION FACTOR 1-ALPHA (FRAGMENT).
 GN TEF1.

OS Paranophrys carnivora.
 OC Eukaryota; Alveolata; Ciliophora; Oligohymenophorea; Scuticociliatida;
 OC Paranophryidae; Paranophrys.
 OX NCBI_TaxID=85900;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE-99152616; PubMed=10028290;
 RA Morella D., Le Guyader H., Philippe H.;
 RT "Unusually high evolutionary rate of the elongation factor 1 alpha
 RT genes from the Ciliophora and its impact on the phylogeny of
 RT eukaryotes.";
 RL Mol. Biol. Evol. 16:234-245(1999).
 CC -1- SIMILARITY: TO GTP-BINDING ELONGATION FACTOR FAMILY.
 DR EMBL: AF056103; AAD03258.1; -
 DR HSSP: Q01698; 1PTT.
 DR InterPro: IPR000795; GTP_EFTU.
 DR Pfam: PF00009; GTP_EFTU; 1.
 DR PRINTS: PR00315; ELONGATINCT.
 DR PROSITE: PS00301; EFACOR_GTP; 1.
 KW Elongation factor; GTP-binding; Protein biosynthesis.
 FT NON_TER 1
 FT NON_TER 408
 SQ SEQUENCE 408 AA; 45066 MW; 384973BB7F3F5FAL CRC64;

Query Match 66.1%; Score 39; DB 5; Length 408;
 Best Local Similarity 77.8%; Pred. No. 50;
 Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY 1 HEAPEAP 9
 Db 269 HESLEAP 277

RESULT 11
 ID O9MAY3 PRELIMINARY; PRT; 417 AA.
 AC O9MAY3;
 DT 01-MAY-2000 (TREMBLrel. 13, Created)
 DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
 DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
 DE CG12497 PROTEIN.
 GN EG:BACR25B3.2 OR CG12497.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
 OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 OX NCBI_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=BERKELEY;
 RX MEDLINE=20196006; PubMed=10731132;
 RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
 RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,
 RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
 RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
 RA Brandon R.C., Rogers Y.H.C., Blaise R.G., Champagne M., Pfeiffer B.D.,
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Milos G.L.G.,
 RA Abriil J.F., Adiyanti A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,
 RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
 RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
 RA Borokova D., Botchan M.R., Bouck J., Brokstein P., Brotlier P.,
 RA Burtis K.C., Busan D.A., Butler H., Cadieu E., Center A., Chandra I.,
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Pablos K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
 RA Durbin K.J., Evangelista C.C., Ferraz C., Ferriera S., Fleischmann W.,
 RA Foster C., Gabrielian A.E., Gary N.S., Gelbart W.M., Glasser K.,
 RA Glodek A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
 RA Hostali D., Houston K.A., Howland J.J., Wei M.-H., Ibeigwan C.,
 RA Jostali D., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,

RA Lasko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
 RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Mostreli A.,
 RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
 RA Nelson D.R., Nelson K.A., Nixon K., Nuskern D.R., Pacle J.M.,
 RA Palazolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
 RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
 RA Slier E., Spralling A.C., Stapleton M., Strong R., Sun E.,
 RA Svirskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
 RA Wang Z.-Y., Wasserman D.A., Weinstein G.M., Weissbach J.,
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
 RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
 RT "The genome sequence of *Drosophila melanogaster*.";
 RL Science 287:2185-2195(2000).
 CC -1- SIMILARITY: TO LOW DENSITY LIPOPROTEIN (LDL) RECEPTOR CLASS A
 (LDLRA) DOMAIN.
 DR EMBL: AE003424; AAF45787.1; -
 DR HSSP: Q07954; 1CR8.
 DR FlyBase: FBgn0040379; EG:BACR25B3.2.
 DR InterPro: IPR002172; LDL_recept_A.
 DR Pfam: PF00057; LDL_recept_a; 1.
 DR PRINTS: PR00261; LDLRECEPTOR.
 DR SMART: SM00192; LDLA; 2.
 DR PROSITE: PS01209; LDLRA_1; 1.
 DR PROSITE: PS50068; LDLRA_2; 2.
 KW Glycoprotein.
 SQ SEQUENCE 417 AA; 48624 MW; 2F1CA7F440D1DD01 CRC64;

Query Match 66.1%; Score 39; DB 5; Length 417;
 Best Local Similarity 87.5%; Pred. No. 51;
 Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 2 EAEPEAP 9
 Db 382 ESEPEAP 389

RESULT 12
 ID O9NEF8 PRELIMINARY; PRT; 435 AA.
 AC O9NEF8;
 DT 01-OCT-2000 (TREMBLrel. 15, Created)
 DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)
 DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
 DE EG:BACR25B3.2 OR CG12497.
 GN EG:BACR25B3.2 OR CG12497.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
 OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 OX NCBI_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Murphy L., Harris D., Bartell B.;
 RT "Sequencing the distal X chromosome of *Drosophila melanogaster*.";
 RL Submitted (FEB-2000) to the EMBL/Genbank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RA Benos P.;
 RT Submitted (FEB-2000) to the EMBL/Genbank/DBJ databases.
 CC -1- SIMILARITY: TO LOW DENSITY LIPOPROTEIN (LDL) RECEPTOR CLASS A
 (LDLRA) DOMAIN.
 DR EMBL: AL138972; CAB72287.1; -
 DR FlyBase: FBgn0040379; EG:BACR25B3.2.
 DR InterPro: IPR002172; LDL_recept_A.
 DR Pfam: PF00057; LDL_recept_a; 1.
 DR PRINTS: PR00261; LDLRECEPTOR.
 DR SMART: SM00192; LDLA; 2.
 DR PROSITE: PS01209; LDLRA_1; 1.

DR PROSITE: PS50068; LDLRA_2; 2.
 KM Glycoprotein.
 SQ SEQUENCE 435 AA; 50714 MW; 19D931A6521C3EAI CRC64;

Query Match
 Best Local Similarity 66.1%; Score 39; DB 5; Length 435;
 Matches 7; Conservativity 1; Mismatches 0; Indels 0; Caps 0;

OY 2 EAEPEAP 9
 Db 400 ESEPEAP 407

RESULT 13

Q21489 PRELIMINARY; PRT; 848 AA.

AC Q21489; 01-NOV-1996 (TREMblrel. 01, Created)

DT 01-NOV-1996 (TREMblrel. 01, Last sequence update)

DE 01-JUN-2001 (TREMblrel. 17, Last annotation update)

GN M03C11.2. PROTEIN.

OS Caenorhabditis elegans.

OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;

NCBI_TaxID=6239;

RN [1]

RP SEQUENCE FROM N.A.

RL Submitted (APR-1995) to the EMBL/GenBank/DBJ databases.

RA MEDLINE=94150718; PubMed=7906398;

RA Wilson R., Ainscough R., Anderson K., Baynes C., Berks M.,

RA Bonfield J., Burton J., Connell M., Copsey T., Cooper J., Coulson A.,

RA Craxton M., Dear S., Du Z., Durbin R., Favello A., Fulton L.,

RA Giermer A., Green P., Hawkins T., Hillier L., Jier M., Johnston L.,

RA Jones M., Kerstan J., Kirsten J., Laister N., Latreille P.,

RA Lightning J., Lloyd C., Murray A., Mortimore B., O'Callaghan M.,

RA Parsons J., Percy C., Rinken L., Roopra A., Saunders D., Shomkneen R.,

RA Smalton N., Smith A., Sonhammer E., Staden R., Sulston J.,

RA Waterson A., Watson K., Vaudin M., Vaughan K., Waterson R.,

RA Watson A., Watson K., Vaudin M., Vaughan K., Waterson R.,

RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.

RT elegans";

RL Nature 368:32-38(1994).

DR EMBL: Z49128; CA88959.1; -

DR InterPro: IPR001410; DEAD.

DR SMART: SM00488; DEXDc2; 1.

DR SMART: SM00491; HELICc2; 1.

SQ SEQUENCE 848 AA; 96811 MW; 4D8574A99F63734B CRC64;

Query Match
 Best Local Similarity 77.8%; Score 39; DB 5; Length 848;
 Matches 7; Conservativity 1; Mismatches 1; Indels 0; Gaps 0;

OY 2 EAEPEAP 10
 Db 506 EPEPEAPL 514

RESULT 14

Q9W475 PRELIMINARY; PRT; 1192 AA.

AC Q9W475; 01-MAY-2000 (TREMblrel. 13, Created)

DT 01-MAY-2000 (TREMblrel. 13, Last sequence update)

DT 01-JUN-2001 (TREMblrel. 17, Last annotation update)

DE CG3108 PROTEIN.

GN CG3108.

OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
 OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.

NCBI_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=BERKELEY.
 RX MEDLINE=20196006; PubMed=10731132;

RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,

RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,

RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,

RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,

RA Brandon R.C., Rogers Y.-H.C., Blazer R.G., Champe M., Pfeiffer B.D.,

RA Man K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Mikos G.L.G.,

RA Adair J.F., Agbayani A., An H.-J., Andrews-Plannkoch C., Baldwin D.,

RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,

RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,

RA Borkova D., Botchan M.R., Bouck J., Brokstein P., Brothier P.,

RA Burtis K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,

RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,

RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,

RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,

RA Durbin K.J., Evangelista C.C., Ferraz C., Ferlita S., Fleischmann W.,

RA Foster C., Gabriellian A.E., Garg N.S., Gehlert W.M., Glasser K.,

RA Glodok A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,

RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,

RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwam C.,

RA Jalali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,

RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,

RA Lasko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,

RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,

RA Merkulov G., Mishina N.V., Mobarry C., Morris J., Moshrefi A.,

RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,

RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacle J.M.,

RA Palazolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,

RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,

RA Shue B.C., Siden-Klamos I., Simpson M., Skupski M.P., Smith T.,

RA Spter E., Spradling A.C., Stapleton M., Strong R., Sun E.,

RA Sytkas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,

RA Williams S.M., Woodage D.A., Watson K.C., Wu D., Yang S., Yao Q.A.,

RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,

RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
 RT "The genome sequence of Drosophila melanogaster";
 RL Science 287:2185-2195(2000).

DR EMBL: AE003435; AAF46083.1; -

DR HSSP: P00730; 2CTC.

DR FlyBase: FBgn0029807; CG3108.

DR InterPro: IPR000834; Zn-carboxypept.

DR Pfam: PF00246; Zn-carboxypept; 1.

DR PRINTS: PR00765; CRBOXYPTASEA.

DR PROSITE: PS00132; CARBOXYPEPT_ZN_1; 1.

SQ SEQUENCE 1192 AA; 132088 MW; 72408D5C5D3D718E CRC64;

Query Match
 Best Local Similarity 66.1%; Score 39; DB 5; Length 1192;
 Matches 7; Conservativity 1; Mismatches 0; Indels 0; Gaps 0;

OY 2 EAEPEAP 9
 Db 330 EAEPEAP 337

RESULT 15

Q08547 PRELIMINARY; PRT; 1277 AA.

AC Q08547; 01-NOV-1996 (TREMblrel. 01, Created)

DT 01-NOV-1996 (TREMblrel. 01, Last sequence update)

DT 01-JUN-2001 (TREMblrel. 17, Last annotation update)

GN CG3108.

TRANS-ACTING TRANSCRIPTIONAL PROTEIN ICP4
 (IMMEDIATE-EARLY PROTEIN IE62).
 62. Cercopithecine herpesvirus 9 (simian varicella virus).
 OS Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
 OC Alphaherpesvirinae; Varicellovirus.
 OX NCBI_TaxID=35246;
 [1]
 RN SEQUENCE FROM N.A.
 RP MEDLINE=94023599; PubMed=8212583;
 RX Clarke P., Brunschwig A., Gildea D.H.;
 RA "DNA sequence of a simian varicella virus gene that encodes a
 RT homologue of varicella zoster virus IE62 and herpes simplex virus
 RT ICP4."
 RL Virology 197:45-52(1993).
 CC -1- FUNCTION: THIS IE PROTEIN IS A MULTIFUNCTIONAL PROTEIN CAPABLE OF
 CC MIGRATING TO THE NUCLEUS, BINDING TO DNA, TRANS-ACTIVATING OTHER
 CC VIRAL GENES, AND AUTOREGULATING ITS OWN SYNTHESIS.
 CC -1- SUBCELLULAR LOCATION: NUCLEUS OF INFECTED CELLS.
 CC -1- PTM: A LONG STRETCH OF SERINE RESIDUES MAY BE A MAJOR SITE OF
 CC PHOSPHORYLATION
 CC -1- SIMILARITY: BELONGS TO THE ICP4/IE140/IE180 FAMILY OF PROTEINS.
 CC EMBL: L20759; AAA03549.1; -;
 DR InterPro: IPR000923; Copper_blue1.
 DR PROSITE: PS00196; CQPPER_BLUE; UNKNOWN_1.
 DR Early protein; Transcription regulation; Trans-acting factor;
 KW DNA-binding; Nuclear protein.
 SO SEQUENCE 1277 AA; 136978 MW; BB92AA8C4DCBD9D CRC64;

Query Match 66.1%; Score 39; DB 12; Length 1277;
 Best local similarity 70.0%; Pred. No. 1.6e+02;
 Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 EAEPEAPEIM 11
 |:|||||:
 Db 1106 EPPEAPEPII 1115

Search completed: January 7, 2002, 08:47:23
 Job time: 411 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw.model

Run on: January 4, 2002, 08:41:02 ; Search time 18.1 seconds
(without alignments)

22.282 Million cell updates/sec

Title: US-09-444-281-27

Sequence: 1 HEAEPFAPEDIM 11

Scoring table: BLOSUM62

Gapop 10.0, Gapext 0.5

Searched: 100059 seqs, 3664827 residues

Total number of hits satisfying chosen parameters: 100059

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Database: Listing first 45 summaries

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match length	ID	Description
1	42	71.2	144 1 AP22.APIME	P35581 apis mellif
2	42	71.2	283 1 AP73.APIME	Q06602 apis mellif
3	40	67.8	429 1 CGB1.CRILIO	Q08301 cricetus
4	39	66.1	486 1 HSL1.HUMAN	P14317 homo sapien
5	38	64.4	25 1 NP4.HUMAN	P18078 homo sapien
6	38	64.4	168 1 AP14.APIME	Q06601 apis mellif
7	38	64.4	248 1 GRL1.RAT	Q06605 rattus norv
8	38	64.4	256 1 PRN3.HUMAN	P24158 homo sapien
9	38	64.4	332 1 H630.YEAST	P25619 saccharomyc
10	37	62.7	226 1 DDN1.BOVIN	P80219 bos taurus
11	37	62.7	245 1 MCT1.SHEEP	P80931 ovils aries
12	37	62.7	246 1 GRAB.HUMAN	P10144 h girazyme
13	37	62.7	247 1 TONB.HUMAN	P20185 serralia ma
14	37	62.7	247 1 TONB.SERNA	P18291 rattus norv
15	37	62.7	248 1 NKPI.RAT	O46683 ovils aries
16	37	62.7	251 1 MCT3.SHEEP	O49942 archaeoglob
17	37	62.7	269 1 VA53.ARCFU	P29940 bacteroides
18	37	62.7	376 1 XYNA.BACOV	P03327 synechococc
19	37	62.7	483 1 PRR.STYNE	P36678 caenorhabdi
20	37	62.7	601 1 PAGT.CAEEL	Q10122 escherichia
21	36	61.0	182 1 YSM2.CAEEL	P02929 escherichia
22	36	61.0	239 1 TONB.ECOLI	P25945 salmonella
23	36	61.0	242 1 TONB.SALTY	O05740 yersinia en
24	36	61.0	255 1 TONB.YEREN	O22712 arbidolipais
25	36	61.0	277 1 PS12.AKATH	Q15759 homo sapien
26	36	61.0	372 1 NK11.HUMAN	P30277 rattus norv
27	36	61.0	423 1 CGB1.RAT	O70324 mus musculu
28	36	61.0	565 1 MCT8.MOUSE	Q16760 homo sapien
29	36	61.0	1195 1 KDGD.HUMAN	P33692 escherichia
30	35	59.3	171 1 ZUR.ECOLI	P54000 saccharomyc
31	35	59.3	292 1 SUB1.YEAST	P45703 bacillus st
32	35	59.3	330 1 XYN2.BACST	P14635 homo sapien
33	35	59.3	433 1 CGB1.HUMAN	

34	35	59.3	465 1 G3BP.MOUSE	P97855 mus musculu
35	35	59.3	466 1 G3BP.HUMAN	Q13283 homo sapien
36	35	59.3	474 1 SY65.DROME	P21521 drosophila
37	35	59.3	486 1 HSL1.MOUSE	P49710 mus musculu
38	35	59.3	613 1 MCT8.HUMAN	P36021 homo sapien
39	35	59.3	754 1 RAD4.YEAST	P14736 saccharomyc
40	35	59.3	788 1 TRS1.HCWA	P09695 human cytom
41	35	59.3	2749 1 IP3R.MOUSE	P18181 mus musculu
42	35	59.3	2749 1 IP3R.RAT	P29994 rattus norv
43	34	57.6	202 1 B3G1.MOUSE	Q9cw73 m galactosy
44	34	57.6	302 1 RS3.HALHA	P15009 halobacteri
45	34	57.6	334 1 B3G1.RAT	O35789 r galactosy

ALIGNMENTS

RESULT	ID	AP22.APIME	STANDARD:	PRT:	144 AA.
AC	P35581	P11525; P11526;			
DT	01-OCT-1989	(Rel. 12, Created)			
DT	01-JUN-1994	(Rel. 29, Last sequence update)			
DT	01-JUN-1994	(Rel. 29, Last annotation update)			
DE	APIDAECIN PRECURSOR, TYPE 22.				
OS	Apis mellifera (Honeybee).				
OC	Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;				
OC	Pterygota; Neoptera; Endopterygota; Hymenoptera; Apocrita; Aculeata;				
OX	Apidae; Apidae; Apis.				
NC	NCBI_TaxID=7460;				
RM	[1]				
RP	SEQUENCE FROM N.A.				
RX	MEDLINE=93223697; PubMed=8467807;				
RA	Casteels-Josson K., Capaci T., Casteels P., Tempst P.;				
RT	"Apidacsin multipetide precursor structure: a putative mechanism for				
RT	amplification of the insect antibacterial response.";				
RL	EMBO J. 12:1569-1578(1993).				
RN	[2]				
RP	SEQUENCE (APIDAECIN IA/IB).				
RC	TISUP-Hemolymph;				
RX	MEDLINE=90005446; PubMed=2676519;				
RA	Casteels P., Ampe C., Jacobs F., Vaecq M., Tempst P.;				
RT	"Apidacsin: antibacterial peptides from honeybees.";				
RL	EMBO J. 8:2387-2391(1989).				
CC	-I- FUNCTION: APIDAECIN HAVE BACTERICIDAL ACTIVITY; PREDOMINANTLY				
CC	AGAINST GRAM-NEGATIVE BACTERIA. THEY SEEM TO INTERFERE WITH CELL				
CC	PROLIFERATION.				
CC	-----				
CC	This SWISS-PROT entry is copyright. It is produced through a collaboration				
CC	between the Swiss Institute of Bioinformatics and the EMBL outstation -				
CC	the European Bioinformatics Institute. There are no restrictions on its				
CC	use by non-profit institutions as long as its content is in no way				
CC	modified and this statement is not removed. Usage by and for commercial				
CC	entities requires a license agreement (see http://www.isb-sib.ch/announce/				
CC	or send an email to license@sib-sib.ch).				
CC	-----				
DR	EMBL; X72576; CAAS1168.1; -				
DR	PIR; S05383; S05383.				
DR	PIR; S06675; S06675.				
DR	PIR; S35331; S35331.				
DR	InterPro: IPR001979; Apidacsin.				
DR	Pfam: PF00807; Apidacsin; 4.				
KW	Insect immunity; Antibiotic; Hemolymph; Signal; Multigene family;				
KW	Cleavage on pair of basic residues; Repeat.				
FT	SIGNAL	1	19	POTENTIAL.	
FT	PROPEP	35	42		
FT	PEPTIDE	43	60	APIDAECIN IB.	
FT	PROPEP	63	70		
FT	PEPTIDE	71	88	APIDAECIN IB.	
FT	PROPEP	91	98		
FT	PEPTIDE	99	116	APIDAECIN IB.	
FT	PROPEP	119	126		
FT	PEPTIDE	127	144	APIDAECIN IA.	

SQ SEQUENCE 144 AA: 16539 MW: 6FA1AD74CB71108D CRC64;

Query Match 71.2%; Score 42; DB 1; Length 144;

Best Local Similarity 100.0%; Pred. No. 1.4;

Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 EAEPAEP 9
11111111
DB 35 EAEPAEP 42

RESULT 2
ID AP73 API ME STANDARD: PRT: 283 AA.

AC 006602: P11525; P11526;
DT 01-OCT-1989 (Rel. 12, Created)
DT 01-JUN-1994 (Rel. 29, Last sequence update)
DT 01-JUN-1994 (Rel. 29, Last annotation update)
DE API DAECIN PRECURSOR, TYPE 73 (FRAGMENT).

OS Apis mellifera (Honeybee).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Hymenoptera; Apocrita; Aculeata;
OC Apoidea; Apidae; Apis.
OX NCBI_TaxID=7460;

RP SEQUENCE FROM N.A.
RX MEDLINE=93223697; PubMed=8467807;
RA Casteels-Josson K., Capaci T., Casteels P., Tempst P.;
RT "Apidaecin multipetide precursor structure: a putative mechanism for
amplification of the insect antibacterial response.";
RL EMBO J. 12:1569-1578(1993).

RN [12]
RP SEQUENCE OF API DAECIN IA/IB.
RC TISSUE=Hemolymph;
RX MEDLINE=90005446; PubMed=2676519;
RA Casteels P., Ampe C., Jacobs F., Vaeck M., Tempst P.;
RT "Apidaecins: antibacterial peptides from honeybees.";
RL EMBO J. 8:2387-2391(1989).
CC -1- FUNCTION: API DAECIN HAVE BACTERICIDAL ACTIVITY: PREDOMINANTLY
AGAINST GRAM-NEGATIVE BACTERIA. THEY SEEM TO INTERFERE WITH CELL
PROLIFERATION.

CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@sib-sib.ch).

CC EMBL: X72577; CAA51169.1; -
DR PIR: S05383; S05383.
DR PIR: S06675; S06675.
DR PIR: S35332; S35332.

DR InterPro: IPR001979; Apidaecin.
DR Pfam: PF00807; Apidaecin.9.
DR ProDom: PD153432; Apidaecin.2.
KW Insect immunity; Antibiotic; Hemolymph; Signal; Multigene family;
KW Cleavage on pair of basic residues; Repeat.

FT SIGNAL 1 18 POTENTIAL.
FT PROPEP 34 41
FT PROPEP 42 59 API DAECIN IB.
FT PROPEP 62 69
FT PROPEP 70 87 *API DAECIN IB.
FT PROPEP 90 97
FT PROPEP 98 115 API DAECIN.
FT PROPEP 118 135
FT PROPEP 126 143 API DAECIN IB.
FT PROPEP 146 153
FT PROPEP 154 171 API DAECIN.
FT *PEPTIDE

FT PROPEP 174 182
FT PEPTIDE 183 199 API DAECIN IB.
FT PROPEP 202 209
FT PEPTIDE 210 227 API DAECIN IB.
FT PROPEP 230 237
FT PEPTIDE 238 255 API DAECIN IB.
FT PROPEP 258 265
FT PEPTIDE 266 283
SQ SEQUENCE 283 AA: 32695 MW: 4EA5FEDEC05E142B CRC64;

Query Match 71.2%; Score 42; DB 1; Length 283;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 EAEPAEP 9
11111111
DB 62 EAEPAEP 69

RESULT 3
ID CGBI_CRI LO STANDARD: PRT: 429 AA.

AC 008301;
DT 01-OCT-1994 (Rel. 30, Created)
DT 01-OCT-1994 (Rel. 30, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE G2/MITOTIC-SPECIFIC CYCLIN B1.

GN CCNB1.
OS Crictetus longicaudatus (Long-tailed hamster) (Chinese hamster).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Cricetinae;
OC Crictetus.
OX NCBI_TaxID=10030;

RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=ovary;
RX MEDLINE=94095439; PubMed=8270434;
RA Markiewicz D.A., McKenna W.G., Flick M.B., Maity A., Muschel R.J.;
RT "The effects of radiation on the expression of a newly cloned and
characterized rat cyclin B mRNA.";
RL Int. J. Radiat. Oncol. Biol. Phys. 28:135-144(1994).
CC -1- FUNCTION: ESSENTIAL FOR THE CONTROL OF THE CELL CYCLE AT THE G2/M
(MITOSIS) TRANSITION.

CC -1- SUBUNIT: INTERACTS WITH THE CDC2 PROTEIN KINASE TO FORM A
CC SERINE/THREONINE KINASE HOLOENZYME COMPLEX ALSO KNOWN AS
CC MATURATION PROMOTING FACTOR (MPF). THE CYCLIN SUBUNIT IMPARTS
CC SUBSTRATE SPECIFICITY TO THE COMPLEX.
CC -1- DEVELOPMENTAL STAGE: ACCUMULATES STEADILY DURING G2 AND IS
CC ABRUPTLY DESTROYED AT MITOSIS.
CC -1- SIMILARITY: BELONGS TO THE CYCLIN FAMILY. CYCLIN AB SUBFAMILY.

CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@sib-sib.ch).

CC EMBL: X64588; CAA5876.1; -
DR PIR: S34224; S34224.
DR HSP: P20248; IJSP.
DR InterPro: IPR000553; Cyclin.
DR Pfam: PF00134; cyclin.1.
DR SMART: SM00385; CYCLIN; 2.
DR PROSITE: PS00292; CYCLINS; 1.
KW Cyclin; Cell cycle; Cell division; Mitosis.
KW DOMAIN 51 86 LYS-RICH.
SQ SEQUENCE 429 AA: 48062 MW: 6E0BAE7511A678B7 CRC64;

Query Match 67.8%; Score 40; DB 1; Length 429;

Best Local Similarity 70.0%; Pred. No. 9.6;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

Oy 2 HAEPEAPIM 11
11111111
Db 98 EPEPEPEPM 107

RESULT 4

HSL_HUMAN STANDARD; PRT; 486 AA.
ID HSL_HUMAN
AC P14317
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE HEMATOPOIETIC LINEAGE CELL SPECIFIC PROTEIN (HEMATOPOIETIC CELL-
SPECIFIC LYN SUBSTRATE 1) (LCKBPL1).
GN HCLSI OR HSL.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
NCBI_TaxID=9606;
RN NCBI_TaxID=9606;
RP SEQUENCE FROM N.A.
RX MEDLINE=90067934; PubMed=2587259;
RA Kitamura D., Kaneko H., Miyagoe Y., Ariyasu T., Watanabe T.;
RT "Isolation and characterization of a novel human gene expressed
specifically in the cells of hematopoietic lineage.";
RL Nucleic Acids Res. 17:9367-9379(1989).
RN [2]
RX SEQUENCE OF 97-108; 193-201 AND 240-248.
RP MEDLINE=9631348; PubMed=8713105;
RA Egerton M., Moritz R.L., Druker B., Kelso A., Simpson R.J.;
RT "Identification of the 70kD heat shock cognate protein (Hsc70) and
alpha-actinin-1 as novel phosphotyrosine-containing proteins in T
lymphocytes.";
RL Biochem. Biophys. Res. Commun. 224:666-674(1996).
RN [3]
RP BINDING TO HAX-1 PROTEIN.
RX MEDLINE=97211841; PubMed=9059808;
RA Suzuki Y., Demoliere C., Kitamura D., Takeshita H., Deuschle U.,
Watanabe T.;
RT "HAX-1, a novel intracellular protein, localized on mitochondria,
directly associates with HSL, a substrate of src family tyrosine
kinases.";
RL J. Immunol. 158:2736-2744(1997).
RN [4]
RX FUNCTION: SUBSTRATE OF THE ANTIGEN RECEPTOR-COUPLED TYROSINE
KINASE. PLAYS A ROLE IN ANTIGEN RECEPTOR SIGNALING FOR BOTH
CLONAL EXPANSION AND DELETION IN LYMPHOID CELLS. DIRECTLY
ASSOCIATES WITH HAX-1, THROUGH BINDING TO ITS C-TERMINAL REGION.
MAY ALSO BE INVOLVED IN THE REGULATION OF GENE EXPRESSION.
CC -1 SUBUNIT: ASSOCIATES WITH THE SH2 AND SH3 DOMAINS OF LCK.
CC -1 SUBCELLULAR LOCATION: MITOCHONDRIAL (PROBABLE).
CC -1 TISSUE SPECIFICITY: EXPRESSED ONLY IN TISSUES AND CELLS OF
HEMATOPOIETIC ORIGIN.
CC -1 DEVELOPMENTAL STAGE: EXPRESSED IN EARLY STAGE OF MYELOID AND
ERYTHROID DIFFERENTIATION.
CC -1 PTM: PHOSPHORYLATED BY LYN RAPIDLY AFTER CROSSLINKING OF SURFACE
TCR ON B CELLS.
CC -1 SIMILARITY: TO CHICKEN P80/85 PROTEINS (CONTACTIN).
CC -1 SIMILARITY: CONTAINS 1 SH3 DOMAIN.
CC THIS SWISS-PROT entry is copyrighted. It is produced through a collaboration
between the Swiss Institute of Bioinformatics and the EMBL outstation -
the European Bioinformatics Institute. There are no restrictions on its
use by non-profit institutions as long as its content is in no way
modified and this statement is not removed. Usage by and for commercial
entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
or send an email to license@sib-sib.ch).
CC EMBL, X16663; CA34651.1; -;
DR PIR: S07633; S07633.
DR HSSP: P29355; 1SEM.

DR MM: 601306; -;
DR InterPro: IPR003134; HSL_rep.
DR InterPro: IPR001452; SH3.
DR Pfam: PF02218; HSL_rep; 4.
DR Pfam: PF00018; SH3; 1.
DR PRINTS: PR00452; SH3DOMAIN.
DR SMART: SM00326; SH3; 1.
DR PROSITE: PS50002; SH3; 1.
KW Repeat: SH3 domain; Phosphorylation.
FT DOMAIN 27 66 INVOLVED IN HAX-1 BINDING.
FT DOMAIN 81 214 3.5 X 37 AA TANDEN REPEATS.
FT REPEAT 81 116 1.
FT REPEAT 117 153 2.
FT REPEAT 154 190 3.
FT REPEAT 191 214 4 (INCOMPLETE).
FT DOMAIN 428 486 SH3.
FT CONFLICT 241 242 KF -> PK (IN REF. 2).
SQ SEQUENCE 486 AA; 53998 MW; 61AE637157DF5DF2 CRC64;

Query Match 66.1%; Score 39; DB 1; Length 486;
Best Local Similarity 77.8%; Pred. No. 16;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Oy 1 HAEPEAPEP 9
11111111
Db 360 YEAPPEPEP 368

RESULT 5

ID NP4_HUMAN STANDARD; PRT; 25 AA.
AC P18078;
DT 01-NOV-1990 (Rel. 16, Created)
DT 01-NOV-1990 (Rel. 16, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE NEUTROPHIL PROTEINASE 4 (EC 3.4.21.-) (NP-4) (FRAGMENT).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
NCBI_TaxID=9606;
RN NCBI_TaxID=9606;
RP SEQUENCE.
RX MEDLINE=91025622; PubMed=2121162;
RA Ohlsson K., Linder C., Rosengren M.;
RT "Monoclonal antibodies specific for neutrophil proteinase 4.
Production and use for isolation of the enzyme.";
RL Biol. Chem. Hoppe-Seyler 371:549-555(1990).
CC -1 TISSUE SPECIFICITY: NEUTROPHILS.
CC -1 SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
TRYPSIN FAMILY.
DR HSSP: P24158; 1FUJ.
DR MEROPS: S01.134; -;
DR InterPro: IPR001254; Trypsin.
DR Pfam: PF00089; Trypsin; 1.
DR PROSITE: PS50240; TRYPSIN_DOM; PARTIAL.
DR PROSITE: PS00134; TRYPSIN_HIS; PARTIAL.
DR PROSITE: PS00135; TRYPSIN_SER; PARTIAL.
KW Hydrolase; Serine protease; Glycoprotein.
FT NON_TER 25
SQ SEQUENCE 25 AA; 2606 MW; B1CB2038274575B0 CRC64;

Query Match 64.4%; Score 38; DB 1; Length 25;
Best Local Similarity 54.5%; Pred. No. 1.2;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

Oy 1 HAEPEAPEPIM 11
11111111
Db 5 HEADPHSRPYM 15

RESULT 6

AP14_APIME STANDARD; PRT: 168 AA.
 ID AP14_APIME STANDARD; PRT: 168 AA.
 AC 006601; P11525; P11526; P11527;
 DT 01-OCT-1989 (Rel. 12, Created)
 DT 01-JUN-1994 (Rel. 29, Last sequence update)
 DT 01-JUN-1994 (Rel. 29, Last annotation update)
 DE APIDAEIN PRECURSOR, TYPE 14.
 GN APID14.
 OS Apis mellifera (Honeybee).
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
 OC Pterygota; Neoptera; Endopterygota; Hymenoptera; Apoecrita; Aculeata;
 OC Apoidea; Apidae; Apis.
 OX NCBI_TaxID=7460;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC MEDLINE=93223697; PubMed=8467807;
 RX Castels J., Jossion K., Capact T., Castels P., Tempst P.;
 RA "Apidaecin multipeptide precursor structure: a putative mechanism for
 RT amplification of the insect antibacterial response.";
 RL EMBO J. 12:1569-1578(1993).
 RN [2]
 RP SEQUENCE OF APIDAEIN IA/IB/II.
 RC TISSUE=Hemolymph;
 RX MEDLINE=90005446; PubMed=2676519;
 RA Castels P., Ampe C., Jacobs F., Vaecck M., Tempst P.;
 RT "Apidaecins: antibacterial peptides from honeybees.";
 RL EMBO J. 8:2387-2391(1989).
 CC -1- FUNCTION: APIDAEIN HAS BACTERICIDAL ACTIVITY; PREDOMINANTLY
 CC AGAINST GRAM-NEGATIVE BACTERIA. THEY SEEM TO INTERFERE WITH CELL
 CC PROPAGATION.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL: X72575; CAA51167.1; -
 DR PIR: S05383; S05383.
 DR PIR: S06675; S06675.
 DR PIR: S06676; S06676.
 DR PIR: S35330; S35330.
 DR InterPro: IPR001979; Apidaecin.
 DR Pfam: PF00807; Apidaecin. 5.
 DR ProDom: PD153432; Apidaecin. 1.
 KM Insect immunity; Antibiotic; Hemolymph; Signal; Multigene family;
 KM Cleavage on pair of basic residues; Repeat.
 KM SIGNAL 1
 FT 19 POTENTIAL.
 FT PROPEP 35 42 APIDAEIN II.
 FT PEPTIDE 43 60 APIDAEIN II.
 FT ACT_SITE 109 109 CHARGE RELAY SYSTEM (BY SIMILARITY).
 FT DISULFID 204 204 CHARGE RELAY SYSTEM (BY SIMILARITY).
 FT DISULFID 143 210 BY SIMILARITY.
 FT DISULFID 174 189 BY SIMILARITY.
 FT CARBOHYD 72 72 N-LINKED (GLCNAC...) (POTENTIAL).
 SQ SEQUENCE 168 AA; 19380 MW; 594B931254C04A37 CRC64;
 QY 2 EAPEEAP 9
 DB 35 EAKPEAP 42
 RESULT 7
 Query Match 64.4%; Score 38; DB 1; Length 168;
 Best Local Similarity 87.5%; Pred. No. 8.3;
 Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

GR1L_RAT STANDARD; PRT: 248 AA.
 ID GR1L_RAT STANDARD; PRT: 248 AA.
 AC 006605;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 01-OCT-1996 (Rel. 34, Last annotation update)
 DE GRANZYME-LIKE PROTEIN I PRECURSOR (EC 3.4.21.-).
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OX NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Ductum;
 RX MEDLINE=93285341; PubMed=8508925;
 RA Amerik A.Y., Jarovoi S.V., Grigorenko V.G., Antonov V.K.;
 RT "Identification, sequence analysis, and characterization of cDNA
 RT clones encoding two granzyme-like serine proteinases from rat
 RT duodenum.";
 RL FEBS Lett. 324:226-230(1993).
 CC -1- FUNCTION: THIS ENZYME IS NECESSARY FOR TARGET CELL LYSIS IN
 CC CELL-MEDIATED IMMUNE RESPONSES.
 CC -1- TISSUE SPECIFICITY: DUODENUM.
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
 CC TRYPSIN FAMILY. STRONGEST TO GRANZYMES AND TO MAST CELL PROTEASES.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL: X66693; CAA47235.1; -
 DR HSP: P04187; 2CPL.
 DR MEROPS: S01.136; -
 DR InterPro: IPR001314; Chymotrypsin.
 DR InterPro: IPR001254; Trypsin.
 DR Pfam: PF00089; trypsin. 1.
 DR PRINTS: PR00722; CHYMOTRYPSIN.
 DR SMART: SM00020; TRYP-Spec. 1.
 DR PROSITE: PS50240; TRYPSIN_DOM. 1.
 DR PROSITE: PS00134; TRYPSIN_HIS. 1.
 DR PROSITE: PS00135; TRYPSIN_SER. 1.
 KM Hydrolyase; Serine protease; Zymogen; Signal; Glycoprotein.
 FT SIGNAL 1 18 BY SIMILARITY.
 FT PROPEP 19 20 ACTIVATION PEPTIDE.
 FT CHAIN 21 248 GRANZYME-LIKE PROTEIN I.
 FT ACT_SITE 65 65 CHARGE RELAY SYSTEM (BY SIMILARITY).
 FT ACT_SITE 109 109 CHARGE RELAY SYSTEM (BY SIMILARITY).
 FT DISULFID 204 204 CHARGE RELAY SYSTEM (BY SIMILARITY).
 FT DISULFID 143 210 BY SIMILARITY.
 FT DISULFID 174 189 BY SIMILARITY.
 FT CARBOHYD 72 72 N-LINKED (GLCNAC...) (POTENTIAL).
 SQ SEQUENCE 248 AA; 27235 MW; 036C81B43A8B972 CRC64;
 QY 1 HEAPEPEPIM 11
 DB 25 HEADPHSRPYM 35
 RESULT 8
 PRN3_HUMAN STANDARD; PRT: 256 AA.
 ID PRN3_HUMAN STANDARD; PRT: 256 AA.
 AC P24158; P15637;
 DT 01-APR-1990 (Rel. 14, Created)

DT 15-DEC-1998 (Rel. 37, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE MYELOBLASTIN PRECURSOR (PC 3.4-21.76) (LEUKOCYTE PROTEINASE 3) (PR-3)
 DE (PR3) (AGP7) (WEGENER'S AUTOANTIGEN) (P29) (C-ANCA ANTIGEN).
 CN PRN3 OR MBN.
 OS Homo sapiens (human).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 NCBI_TaxId=9606;
 [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=92021028; PubMed=1681549;
 RA Labbaye C., Musette P., Cayre Y.E.;
 RT "Wegener autoantigen and myeloblastin are encoded by a single mRNA.";
 RL Proc. Natl. Acad. Sci. U.S.A. 88:9253-9256(1991).
 [2]
 RP SEQUENCE FROM N.A.
 RX Lamedin J.E., McGready P.M., Skowronski E., Adamson A.W.,
 RA Burkhart-Schultz R.J., Gordon L., Kyle A., Ramirez M., Stillwagen S.,
 RA Phan H., Velasco N., Do L., Regala W., Terry A., Garnes J.,
 RA Dangnan L., Poundstone P., Christensen M., Georgescu A., Avila J.,
 RA Liu S., Attix C., Andeise T., Frankheim M., Amico-Keller G.,
 RA Coefield J., Duarte S., Lucas S., Bruce R., Thomas P., Quan G.,
 RA Krommiller B., Arellano A., Montgomery M., Ow D., Nolan M., Trong S.,
 RA Kobayashi A., Olsen A.S., Carrano A.V.;
 RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
 [3]
 RP SEQUENCE OF 2-256 FROM N.A. AND SEQUENCE OF 48-71 AND 156-181.
 RX MEDLINE=91079774; PubMed=2258701;
 RA Campanelli D., Melchior M., Fu Y., Nakata M., Shuman H., Nathan C.,
 RA Gabay J.E.;
 RT "Cloning of cDNA for proteinase 3: a serine protease, antibiotic, and
 RT autoantigen from human neutrophils.";
 RL J. Exp. Med. 172:1709-1715(1990).
 [4]
 RP SEQUENCE OF 1-20 AND 22-256 FROM N.A.
 RX MEDLINE=92390417; PubMed=1518849;
 RA Zimmer M., Medcalf R.L., Fink T.M., Maltmann C., Lichter P.,
 RA Jenne D.E.;
 RT "Three human elastase-like genes coordinately expressed in the
 RT myelomonocyte lineage are organized as a single genetic locus on
 RT 19pter.";
 RL Proc. Natl. Acad. Sci. U.S.A. 89:8215-8219(1992).
 [5]
 RP SEQUENCE OF 42-256 FROM N.A.
 RX MEDLINE=90090622; PubMed=2596267;
 RA Borjes D., Raynal M.-C., Solomon D.H., Darzynkiewicz Z., Cayre Y.E.;
 RT "Down-regulation of a serine protease, myeloblastin, causes growth
 RT arrest and differentiation of promyelocytic leukemia cells.";
 RL Cell 59:959-968(1989).
 [6]
 RP SEQUENCE OF 28-67 AND 228-244.
 RX MEDLINE=91236723; PubMed=2033050;
 RA Rao N.V., Wehner N.G., Marshall B.C., Gray W.R., Gray B.H.,
 RA Hoidal J.R.;
 RT "Characterization of proteinase-3 (PR-3), a neutrophil serine
 RT proteinase. Structural and functional properties.";
 RL J. Biol. Chem. 266:9540-9548(1991).
 [7]
 RP SEQUENCE OF 28-47.
 RX MEDLINE=89315847; PubMed=2501794;
 RA Gabay J.E., Scott R.W., Campanelli D., Griffith J., Wilde C.,
 RA Maria M.N., Seeger M., Nathan C.F.;
 RT "Antibiotic proteins of human polymorphonuclear leukocytes.";
 RL Proc. Natl. Acad. Sci. U.S.A. 86:5610-5614(1989).
 [8]
 RP SEQUENCE OF 28-47 AND 196-219.
 RX MEDLINE=90130450; PubMed=2404977;
 RA Wilde C.G., Snable J.L., Griffith J.E., Scott R.W.;
 RT "Characterization of two azurophilic granule proteases with active-site
 RT homology to neutrophil elastase.";
 RL J. Biol. Chem. 265:2038-2041(1990).
 [9]

RP SEQUENCE OF 28-48, AND IDENTITY OF WEGENER'S AUTOANTIGEN WITH PR-3.
 RX MEDLINE=9032035; PubMed=2377228;
 RA Jenne D.E., Tschopp J., Luedemann J., Utecht B., Gross W.L.;
 RT "Wegener's autoantigen decoded.";
 RL Nature 346:520-520(1990).
 [10]
 RP IDENTITY OF WEGENER'S AUTOANTIGEN WITH PROTEINASE 3.
 RX MEDLINE=91055123; PubMed=2242436;
 RA Gupta S.K., Niles J.L., McCluskey R.T., Annout M.A.;
 RT "Identity of Wegener's autoantigen (p29) with proteinase 3 and
 RT myeloblastin.";
 RL Blood 76:2162-2162(1990).
 [11]
 RP X-RAY CRYSTALLOGRAPHY (2.2 ANGSTROMS).
 RX MEDLINE=96346178; PubMed=8757293;
 RA Fujinaga M., Charnala M.M., Halenbeck R., Kothe K., James M.N.G.;
 RT "The crystal structure of PR3, a neutrophil serine proteinase antigen
 RT of Wegener's granulomatosis antibodies.";
 RL J. Mol. Biol. 261:267-278(1996).
 CC -1- FUNCTION: POLYMORPHONUCLEAR LEUKOCYTE SERINE PROTEASE THAT
 CC DEGRADES ELASTIN, FIBRONECTIN, LAMININ, VITRONECTIN, AND COLLAGEN
 CC TYPES I, III, AND IV (IN VITRO) AND CAUSES EMPHYSEMA WHEN
 CC ADMINISTERED BY TRACHEAL INSUFFLATION TO HAMSTERS.
 CC -1- CATALYTIC ACTIVITY: HYDROLYSIS OF PROTEINS, INCLUDING ELASTIN, BY
 CC PREFERENTIAL CLEAVAGE: ALA-I-XAA > VAL-I-XAA.
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
 CC TRYPSIN FAMILY ELASTASE SUBFAMILY.
 CC -----
 CC THIS SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL: X56132; CAA39598.1; -;
 DR EMBL: AC004799; AAC18958.1; -;
 DR EMBL: M75154; AAA59558.1; -;
 DR EMBL: M96839; AAB58493.1; -;
 DR EMBL: M96836; AAB59493.1; JOINED.
 DR EMBL: M96837; AAB59493.1; JOINED.
 DR EMBL: X55668; CAA39203.1; -;
 DR EMBL: M29142; AAA36342.1; -;
 DR EMBL: M96628; AAB59364.1; -;
 DR PIR: A43983; PRH03.
 DR PDB: 1F0J; 11-JUL-96.
 DR MEROPS: S01.134; -;
 DR MIM: 177020; -;
 DR InterPro: IPR001314; Chymotrypsin.
 DR InterPro: IPR001254; Trypsin.
 DR Pfam: PF00089; Trypsin.1.
 DR PRINTS: PR00722; CHYMOTRYPSIN.
 DR SMART: SM00020; TRYP_SPC.1.
 DR PROSITE: PSS0240; TRYPSIN_DOM.1.
 DR PROSITE: PS00134; TRYPSIN_HIS.1.
 DR PROSITE: PS00135; TRYPSIN_SER.1.
 KW Hydroxylase; Serine protease; Glycoprotein; Zymogen; Signal;
 KW 3D-structure.
 FT SIGNAL 1 25
 FT PROPEP 26 27
 FT CHAIN 28 248
 FT PROPEP 249 256
 FT ACT_SITE 71 71
 FT ACT_SITE 118 118
 FT ACT_SITE 203 203
 FT CARBOHYD 129 129
 FT CARBOHYD 174 174
 FT DISULFID 56 72
 FT DISULFID 152 209
 FT DISULFID 182 188
 FT DISULFID 199 224
 FT CONFLICT 2 2
 A -> R (IN REF. 3).

SQ SEQUENCE 226 AA; 25051 MW; A4CAZCE66736CD12 CRC64;
 Query Match 62.7%; Score 37; DB 1; Length 226;
 Best Local Similarity 54.5%; Pred. No. 17;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;
 Oy 1 HEAPEAPEPM 11
 |||:|:|:|
 Db 5 HEAPEHSPYM 15
 RESULT 11
 MCTL_SHEEP STANDARD; PRT: 245 AA.
 ID MCTL_SHEEP
 AC P80931;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE MAST CELL PROTEASE 1A PRECURSOR (EC 3.4.21.-) (SMCP-1A).
 OS Ovis aries (Sheep).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovidae; Caprinae; Ovis.
 OC NCBI_TaxID=9940;
 OX [1]
 RP SEQUENCE FROM N.A. AND SEQUENCE OF 176-185.
 RC TISSUE-Mast cells;
 RX MEDLINE=98343972; PubMed=9677343.
 RA McAliese S.M., Pemberton A.D., McGrath M.E., Huntley J.F.,
 RA Miller H.R.P.;
 RT "Sheep mast-cell proteinases-1 and -3: cDNA cloning, primary
 RT structure and molecular modelling of the enzymes and further studies
 RT on substrate specificity."
 RL Biochem. J. 333:801-809(1998).
 RN [2]
 RP SEQUENCE OF 20-44.
 RA Miller H.R.P., Huntley J.F., Newlands G.F.J.;
 RL (in) Caughey G.H. (eds.);
 RL Mast cell proteases in immunology and biology, pp.203-235, Marcel
 RL Dekker, New York (1995).
 RN [3]
 RP SEQUENCE OF 20-29.
 RC TISSUE-Gastric mucosa;
 RX MEDLINE=97184650; PubMed=9032451.
 RA Pemberton A.D., Huntley J.F., Miller H.R.P.;
 RT "Sheep mast cell proteinase-1: characterization as a member of a new
 RT class of dual-specific ruminant chymases."
 RL Biochem. J. 321:665-670(1997).
 CC -1- FUNCTION: HAS A CHYMOTRYPSIN-LIKE AND TRYPSIN-LIKE ACTIVITY.
 CC -1- SUBCELLULAR LOCATION: SECRETORY GRANULES.
 CC -1- TISSUE SPECIFICITY: MUCOSAL MAST CELLS.
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
 CC TRYPSIN FAMILY. STRONGEST TO OTHER GRANZYMES AND TO MAST CELL
 CC PROTEASES.
 CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL Outstation
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC
 CC EMBL: Y14654; CAA74984.1; -
 CC HSSP: P00763; IDPO.4
 CC DR MEROPS: S01.142; -
 CC DR InterPro: IPR001314; Chymotrypsin.
 CC DR InterPro: IPR001254; Trypsin.
 CC DR Pfam: PF00089; trypsin.1
 CC DR PRINTS: PR00722; CHYMOTRYPSIN.
 CC DR SMART: SM00020; TRYP_SPC.1
 CC PROSITE: PS50240; TRYPSIN_DOM.1.

DR PROSITE: PS00134; TRYPSIN_HIS.1.
 DR PROSITE: PS00135; TRYPSIN_SER.1.
 KM Hydrolyase; Serine protease; Zymogen; Signal.
 FT SIGNAL 1 17
 FT PROPEP 18 19 ACTIVATION PEPTIDE.
 FT CHAIN 20 245 MAST CELL PROTEASE 1A.
 FT ACT_SITE 63 63 CHARGE RELAY SYSTEM (BY SIMILARITY).
 FT ACT_SITE 107 107 CHARGE RELAY SYSTEM (BY SIMILARITY).
 FT ACT_SITE 201 201 CHARGE RELAY SYSTEM (BY SIMILARITY).
 FT DISULFID 48 64 BY SIMILARITY.
 FT DISULFID 141 207 BY SIMILARITY.
 FT DISULFID 172 186 BY SIMILARITY.
 FT CONFLICT 20 20 I -> F (IN REF. 2; AA SEQUENCE).
 SQ SEQUENCE 245 AA; 26877 MW; C362CF3367FFFI80 CRC64;
 Query Match 62.7%; Score 37; DB 1; Length 245;
 Best Local Similarity 54.5%; Pred. No. 18;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;
 Oy 1 HEAPEAPEPM 11
 |||:|:|:|
 Db 24 HEAPEHSPYM 34
 RESULT 12
 GRAH_HUMAN STANDARD; PRT: 246 AA.
 ID GRAH_HUMAN
 AC P20718;
 DT 01-FEB-1991 (Rel. 17, Created)
 DT 01-FEB-1991 (Rel. 17, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE GRANZYME H PRECURSOR (EC 3.4.21.-) (CYTOTOXIC T-LYMPHOCYTE PROTEINASE)
 DE (CATHEPSIN G-LIKE 2) (CTSGL2) (CCP-X) (CYTOTOXIC SERINE PROTEASE-C)
 DE (CSP-C).
 GN GZMH OR CTSGL2 OR CGL2.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 OC NCBI_TaxID=9606;
 OX [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=90298167; PubMed=2193684;
 RA Meier M., Kwong P.C., Fregeau C.J., Atkinson E.A., Burrington M.,
 RA Ehtman N., Sorensen O., Lin C.C., Wilkins J., Bleackley R.C.;
 RT "Cloning of a gene that encodes a new member of the human cytotoxic
 RT cell protease family."
 RL Biochemistry 29:4042-4049(1990).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=91265452; PubMed=2049336;
 RA Haddad P., Jenne D.E., Tschopp J., Clement M.V., Mathieu-Manul D.,
 RA Sasportes M.;
 RT "Structure and evolutionary origin of the human granzyme H gene."
 RL Int. Immunol. 3:57-66(1991).
 RN [3]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=90385483; PubMed=2402757;
 RA Klein J.L., Selvakumar A., Tripathi J.A., Dupont B.;
 RT "Characterization of a novel, human cytotoxic lymphocyte-specific
 RT serine protease cDNA clone (CSP-C)."
 RL Tissue Antigens 35:220-228(1990).
 CC -1- FUNCTION: THIS ENZYME IS PROBABLY NECESSARY FOR TARGET CELL
 CC LYSIS IN CELL-MEDIATED IMMUNE RESPONSES.
 CC -1- SUBCELLULAR LOCATION: CYTOPLASMIC GRANULES OF CYTOLYTIC
 CC T-LYMPHOCYTES.
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
 CC TRYPSIN FAMILY. STRONGEST TO OTHER GRANZYMES AND TO MAST CELL
 CC PROTEASES.
 CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL Outstation
 CC the European Bioinformatics Institute. There are no restrictions on its

CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (see <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).

CC -----
DR EMBL: J02907; AAA76859.1; -
DR EMBL: M57888; AAA03514.1; -
DR EMBL: M36118; AAA03248.1; -
DR EMBL: M72150; AAA74885.1; -
DR PIR: A32692; A32692.
DR HSSP: P04187; 2CP1.
DR MEROPS: S01.147; -.
DR MIM: 116831; -.
DR InterPro: IPR001314; Chymotrypsin.
DR InterPro: IPR001254; Trypsin.
DR Pfam: PF00089; Trypsin_1.
DR PRINTS: PR00722; CHYMOTRYPSIN.
DR SMART: SM00020; TRYP_SPC; 1.
DR PROSITE: PS0240; TRYPSIN_DOM; 1.
DR PROSITE: PS00134; TRYPSIN_HIS; 1.
DR PROSITE: PS00135; TRYPSIN_SER; 1.
KW Hydrolase; Serine protease; Zymogen; Signal; T-cell; Cytolysis.
FT SIGNAL 1 18
FT PROPEP 19 20 ACTIVATION PEPTIDE.
FT CHAIN 21 246 GRANZYME H.
FT ACT_SITE 64 64 CHARGE RELAY SYSTEM (BY SIMILARITY).
FT ACT_SITE 108 108 CHARGE RELAY SYSTEM (BY SIMILARITY).
FT ACT_SITE 202 202 CHARGE RELAY SYSTEM (BY SIMILARITY).
FT DISULFID 49 65 BY SIMILARITY.
FT DISULFID 142 208 BY SIMILARITY.
FT DISULFID 172 187 BY SIMILARITY.
FT CARBOHYD 71 71 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 104 104 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 179 179 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 246 AA: 27315 MW: CA6A87E3DA5F1E71 CRC64;

Query Match 62.7%; Score 37; DB 1; Length 246;
Best Local Similarity 54.5%; Pred. No. 18;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 1 HEAPEAEPTM 11
| | | | |
DB 25 HEAKPHSRPYM 35

RESULT 13
GRAB_HUMAN STANDARD; PRT: 247 AA.
ID GRAB_HUMAN STANDARD; PRT: 247 AA.
AC P10144;
DT 01-MAR-1989 (Rel. 10; Created)
DT 01-MAR-1989 (Rel. 10; Last sequence update)
DT 20-AUG-2001 (Rel. 40; Last annotation update)
DE GRANZYME B PRECURSOR (EC 3.4.21.79) (T-CELL SERINE PROTEASE 1-3F)
DE (CYTOTOXIC T-LYMPHOCYTE PROTEINASE 2) (LYMPHOCYTE PROTEASE) (SECT)
DE (GRANZYME 2) (CATHEPSIN G-LIKE 1) (CTSG1) (CTLA-1) (FRAGMENTIN 2)
DE (HUMAN LYMPHOCYTE PROTEIN) (HLP) (C11).
GN GZMB OR CTLA1 OR GRB OR CGBP OR CCL1.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=87224164; PubMed=2953613;
RA Schmid J., Weissman C.;
RT "Induction of mRNA for a serine protease and a
RT beta-thromboglobulin-like protein in mitogen-stimulated human
RT leukocytes.";
RL J. Immunol. 139:250-256(1987).
RP SEQUENCE FROM N.A.
RX MEDLINE=88198184; PubMed=3258665;

RA Caputo A., Fahey D., Lloyd C., Vozab R., McCalins E., Rowe P.B.;
RT "Structure and differential mechanisms of regulation of expression of
RT a serine esterase gene in activated human T lymphocytes.";
RL J. Biol. Chem. 263:6363-6369(1988).
RN [3]
RP SEQUENCE FROM N.A.
RX MEDLINE=88320548; PubMed=3261871;
RA Trapani J.A., Klein J.L., White P.C., Dupont B.;
RT "Molecular cloning of an inducible serine esterase gene from human
RT cytotoxic lymphocytes.";
RL Proc. Natl. Acad. Sci. U.S.A. 85:6924-6928(1988).
RN [4]
RP SEQUENCE FROM N.A.
RX MEDLINE=89357968; PubMed=2788607;
RA Klein J.L., Shows T.B., Dupont B., Trapani J.A.;
RT "Genomic organization and chromosomal assignment for a serine
RT protease gene (CSPB) expressed by human cytotoxic lymphocytes.";
RL Genomics 5:110-117(1989).
RN [5]
RP SEQUENCE FROM N.A.
RX MEDLINE=90308320; PubMed=2365998;
RA Caputo A., Sauer D.E., Rowe P.B.;
RT "Nucleotide sequence and genomic organization of a human T lymphocyte
RT serine protease gene.";
RL J. Immunol. 145:737-744(1990).
RN [6]
RP SEQUENCE FROM N.A.
RX MEDLINE=90236319; PubMed=2332171;
RA Haddad P., Clement M.V., Bernard O., Larsen C.J., Degos L.,
RA Sasportes M., Mathieu-Mahul D.;
RT "Structural organization of the hCTLA-1 gene encoding human granzyme
RT B.";
RL Gene 87:265-271(1990).
RN [7]
RP SEQUENCE OF 21-40, AND CHARACTERIZATION.
RX MEDLINE=8909866; PubMed=3262682;
RA Hameed A., Lowrey D.M., Lichtenheld M., Podack E.R.;
RT "Characterization of three serine esterases isolated from human IL-2
RT activated killer cells.";
RL J. Immunol. 141:3142-3147(1988).
RN [8]
RP SEQUENCE OF 21-40, AND CHARACTERIZATION.
RX MEDLINE=89035468; PubMed=3263427;
RA Kraehenbuhl O., Rey C., Jenne D.E., Lanzavecchia A., Groscurth P.,
RA Carrel S., Tschopp J.;
RT "Characterization of granzymes A and B isolated from granules of
RT cloned human cytotoxic T lymphocytes.";
RL J. Immunol. 141:3471-3477(1988).
RN [9]
RP SEQUENCE OF 21-38.
RX MEDLINE=91093203; PubMed=1985927;
RA Poe M., Blake J.T., Boulton D.A., Gammon M., Sigal N.H., Wu J.K.,
RA Zweerink H.J.;
RT "Human cytotoxic lymphocyte granzyme B: its purification from
RT granules and the characterization of substrate and inhibitor
RT specificity.";
RL J. Biol. Chem. 266:98-103(1991).
RN [10]
RP FUNCTION: THIS ENZYME IS NECESSARY FOR TARGET CELL LYSIS IN CELL-
RP MEDIATED IMMUNE RESPONSES. IT CLEAVES AFTER ASP. SEEMS TO BE
RP LINKED TO AN ACTIVATION CASCADE OF CASPASES (ASPARTATE-SPECIFIC
RP CYSTEINE PROTEASES) RESPONSIBLE FOR APOPTOSIS EXECUTION. CLEAVES
RP CASPASE-3, -7, -9 AND 10 TO GIVE RISE TO ACTIVE ENZYMES MEDIATING
RP APOPTOSIS.
RN [11]
RP CATALYTIC ACTIVITY: PREFERENTIAL CLEAVAGE: ASP-|-XAA >> ASN-|-XAA
RN [12]
RP SUBCELLULAR LOCATION: CYTOPLASMIC GRANULES OF CYTOLYTIC T-
RN [13]
RP LYMPOCYTES AND NATURAL KILLER CELLS.
RN [14]
RP INDUCTION: BY STAPHYLOCOCCAL ENTEROTOXIN A (SEA) IN PERIPHERAL
RN [15]
RP BLOOD LEUKOCYTES.
RN [16]
RP SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1: ALSO KNOWN AS THE
RN [17]
RP TRYPSIN FAMILY. STRONGEST TO OTHER GRANZYMES AND TO MAST CELL
RN [18]
RP PROTEASES.

CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@sib-sib.ch).

DR EMBL: M17016; AAA36627.1; -
 DR EMBL: J03189; AAA36603.1; -
 DR EMBL: J04071; AAA52118.1; -
 DR EMBL: J03072; AAB59528.1; -
 DR EMBL: M38193; AAB67124.1; -
 DR EMBL: M28879; AAA75490.1; -
 DR PIR: A28659; A28659; -
 DR PIR: A31405; A31405; -
 DR PIR: A32168; A32168; -
 DR PIR: B30525; B30525; -
 DR PIR: B30526; B30526; -
 DR HSSP: P04187; 2CP1; -
 DR MEROPS: S01.010; -
 DR MIM: I23910; -
 DR InterPro: IPR001254; Trypsin.
 DR Pfam: PF00089; trypsin; 1.
 DR SMART: SM00020; TRYP_SPE; 1.
 DR PROSITE: PS50240; TRYP_SIN_DOM; 1.
 DR PROSITE: PS00134; TRYP_SIN_HIS; 1.
 DR PROSITE: PS00135; TRYP_SIN_SER; 1.
 DR K01: Hydrolyase; Serine protease; Zymogen; Signal; T-cell; Cytolysis;
 KW Apoptosis.
 FT SIGNAL 1 18
 FT PROPEP 19 20
 FT CHARIN 21 247
 FT ACT_SITE 64 247
 FT ACT_SITE 108 108
 FT ACT_SITE 203 203
 FT DISULFID 49 65
 FT DISULFID 142 209
 FT DISULFID 173 188
 FT CARBOHYD 71 71
 FT CARBOHYD 104 104
 FT CONFLICT 55 55
 FT CONFLICT 72 72
 FT CONFLICT 94 94
 FT CONFLICT 212 212
 FT CONFLICT 247 AA; 27688 MM; 684FF605D6F2F4FB CRC64;
 SQ SEQUENCE

Query Match 62.7%; Score 37; DB 1; Length 247;
 Best Local Similarity 54.5%; Pred. No. 18;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPIM 11
 |||:|:|
 Db 25 HEAKPHSRPYM 35

RESULT 14
 TONB_SERMA STANDARD; PRT; 247 AA.
 AC P26185;
 DT 01-MAY-1992 (Rel. 22, Created)
 DT 01-MAY-1992 (Rel. 22, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE TONB PROTEIN.
 GN TONB.
 OS Serratia marcescens.
 CC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
 CC Serratia.
 CC NCBI_TaxID=615;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=W225;

RX MEDLINE=92140042; PubMed=1838128;
 RA Gaiser S., Braun V.:
 RT "The tonB gene of Serratia marcescens: sequence, activity and partial
 RT complementation of Escherichia coli tonB mutants.";
 RL Mol. Microbiol. 5:2777-2787 (1991).
 CC -1- FUNCTION: INTERACTS WITH OUTER MEMBRANE RECEPTOR PROTEINS THAT
 CC CARRY OUT HIGH-AFFINITY BINDING AND ENERGY DEPENDENT UPTAKE INTO
 CC THE PERIPLASMIC SPACE OF SPECIFIC SUBSTRATES SUCH AS COBALAMIN,
 CC AND VARIOUS IRON COMPOUNDS (SUCH AS IRON DICTYRATE, ENTEROCHELIN,
 CC AEROBACTIN, ETC.). IN THE ABSENCE OF TONB THESE RECEPTORS BIND
 CC THEIR SUBSTRATES BUT DO NOT CARRY OUT ACTIVE TRANSPORT. TONB ALSO
 CC INTERACTS WITH SOME COLICINS AND IS INVOLVED IN THE ENERGY-
 CC DEPENDENT, IRREVERSIBLE STEPS OF BACTERIOPHAGES PH1-80 AND T1
 CC INFECTION. IT COULD ACT TO TRANSDUCE ENERGY FROM THE CYTOPLASMIC
 CC MEMBRANE TO SPECIFIC ENERGY-REQUIRING PROCESSES IN THE OUTER
 CC MEMBRANE, RESULTING IN THE RELEASE INTO THE PERIPLASM OF LIGANDS
 CC BOUND BY THESE OUTER MEMBRANE PROTEINS (BY SIMILARITY).
 CC -1- SUBUNIT: THE ACCESSORY PROTEINS EXBB AND EXBD SEEM TO FORM A
 CC COMPLEX WITH TONB.
 CC -1- SUBCELLULAR LOCATION: PERIPLASMIC. ANCHORED TO THE CYTOPLASMIC
 CC MEMBRANE VIA ITS N-TERMINAL SIGNAL-LIKE SEQUENCE. SPANS THE
 CC PERIPLASM.
 CC -1- SIMILARITY: BELONGS TO THE TONB FAMILY.

CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@sib-sib.ch).

DR EMBL: X60996; GAA43308.1; -
 DR InterPro: IPR003538; TonB.
 DR Transport; Protein transport; Bacteriocin transport; Inner membrane;
 KW Periplasmic; Transmembrane; Signal-anchor; Repeat; Phage recognition.
 FT DOMAIN 1 10
 FT TRANSMEM 12 35
 FT DOMAIN 36 247
 FT DOMAIN 76 85
 FT DOMAIN 101 110
 FT DOMAIN 247 AA; 27389 MM; 46EE6869EDB864B CRC64;
 SQ SEQUENCE

Query Match 62.7%; Score 37; DB 1; Length 247;
 Best Local Similarity 70.0%; Pred. No. 18;
 Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 2 EAPEPEPEPIM 11
 |||||
 Db 78 EPEPEPEPIV 87

RESULT 15
 NKPL_RAT STANDARD; PRT; 248 AA.
 AC P18291;
 DT 01-NOV-1990 (Rel. 16, Created)
 DT 01-NOV-1990 (Rel. 16, Last sequence update)
 DT 01-OCT-1996 (Rel. 34, Last annotation update)
 DE NATURAL KILLER CELL PROTEASE 1 PRECURSOR (EC 3.4.21.-) (RNKP-1)
 DE (FRAGMENTIN).
 OS Rattus norvegicus (Rat).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 CC NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RP TISSUE=T-cell;
 RX MEDLINE=90171602; PubMed=2307850;
 RA Zunino S.J., Bleackley R.C., Martinez J., Hudig D.:
 RT "RNKP-1, a novel natural killer cell-associated serine protease gene
 RT cloned from RNK-16 cytotoxic lymphocytes.";

RL J. Immunol. 144:2001-2009(1990).
RN [2]
RP SEQUENCE OF 21-53.
RX MEDLINE-92091788; PubMed=1727874;
RA Sayers T.J., Wiltrout T.A., Sowder R., Munger W.L., Smyth M.J.,
RA Henderson L.E.;
RT "Purification of a factor from the granules of a rat natural killer
RT cell line (RNK) that reduces tumor cell growth and changes tumor
RT morphology. Molecular identity with a granule serine protease
RT (RNKP-1).";
RL J. Immunol. 148:292-300(1992).
RN [3]
RP PARTIAL SEQUENCE.
RX MEDLINE-92121838; PubMed=1732416;
RA Shi L., Kraut R.P., Aebersold R., Greenberg A.H.;
RT "A natural killer cell granule protein that induces DNA fragmentation
RT and apoptosis.";
RL J. Exp. Med. 175:553-566(1992).
CC -1- FUNCTION: THIS ENZYME IS PROBABLY NECESSARY FOR TARGET CELL
CC LYSIS IN CELL-MEDIATED IMMUNE RESPONSES.
CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
CC TRYPSIN FAMILY. STRONGEST TO OTHER GRANZYMES AND TO MAST CELL
CC PROTEASES.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-stb.ch/announce/>
CC or send an email to license@isb-stb.ch).
CC -----
DR EMBL: M34097; AAA42055.1; -
DR PIR: A43520; A43520.
DR HSSP: P04187; 2CPI.
DR MEROPS: S01.136; -
DR InterPro: IPR001254; Trypsin.
DR Pfam: PF00089; Trypsin.1.
DR SMART: SM00020; TRYP_SPC; 1.
DR PROSITE: PS50240; TRYPSIN_DOM; 1.
DR PROSITE: PS00134; TRYPSIN_HIS; 1.
DR PROSITE: PS00135; TRYPSIN_SER; 1.
KW Hydrolase; Serine protease; Zymogen; Signal; T-cell; Cytolysis.
KW SIGNAL
FT PROPEP 1 18
FT CHAIN 19 20
FT ACT_SITE 21 248
FT ACT_SITE 65 65
FT ACT_SITE 109 109
FT ACT_SITE 204 204
FT DISULFID 50 66
FT DISULFID 143 210
FT DISULFID 174 189
FT DISULFID 98 98
FT CONFLICT 138 138
SQ SEQUENCE 248 AA; 27326 MW; 6F52089DDACC88C CRC64;

Query Match 62.7%; Score 37; DB 1; Length 248;
Best Local Similarity 54.5%; Pred. No. 18;
Matches 6: Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 1 HEAEPEAEPIIM 11
|||:|:|
DB 25 HEAKPHSRPYM 35

GenCone version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:38:42 ; Search time 27.18 Seconds
(without alignments)
30.829 Million cell updates/sec

Title: US-09-444-281-27
Perfect score: 59
Sequence: 1 HEAPEAPEIM 11

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 219241 seqs, 76174552 residues
Total number of hits satisfying chosen parameters: 219241

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: p1r1:*
2: p1r2:*
3: p1r3:*
4: p1r4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	42	71.2	144	2	S35331 apidaecin 22 precu
2	42	71.2	283	2	S35332 apidaecin 73 precu
3	40	67.8	429	2	S34224 cyclin B - long-ta
4	39	66.1	251	2	T39332 hypothetical prote
5	39	66.1	486	2	S07633 hematopoietic line
6	39	66.1	848	2	T23694 hypothetical prote
7	39	66.1	1593	2	T22028 hypothetical prote
8	38	64.4	150	2	C72724 hypothetical prote
9	38	64.4	168	2	S35330 apidaecin 14 precu
10	38	64.4	248	2	S33755 granzyme-like prot
11	38	64.4	256	1	PRH03 granzyme-3 like prot
12	38	64.4	288	2	T51059 proteinase 3 (EC 3
13	38	64.4	332	2	S31848 hypothetical prote
14	37	62.7	21	2	S69371 heat shock protein
15	37	62.7	125	2	T36367 duodenase - bovine
16	37	62.7	175	2	T05669 hypothetical prote
17	37	62.7	226	2	S69370 duodenase - bovine
18	37	62.7	246	2	A32692 cytochrome P-1 lympo
19	37	62.7	247	2	S18592 tonB protein - Ser
20	37	62.7	248	2	S43259 granzyme-like prot
21	37	62.7	248	2	A43520 natural killer cel
22	37	62.7	251	2	T10262 mast cell serine p
23	37	62.7	265	2	B81138 phosphatidylserine
24	37	62.7	265	2	C81883 probable membrane
25	37	62.7	269	2	E69381 hypothetical prote
26	37	62.7	281	1	A61021 granzyme B (EC 3.4
27	37	62.7	356	2	T00881 probable PCP2-like
28	37	62.7	376	2	S55892 endo-1,4-beta-xyla
29	37	62.7	484	2	S00757 deoxyribodipyrimid

30	37	62.7	612	2	T42243 probable polypepti
31	37	62.7	1280	2	T00365 hypothetical prote
32	37	62.7	2109	2	T33247 hypothetical prote
33	37	62.7	2584	2	T24158 hypothetical prote
34	37	62.7	2606	2	T24157 hypothetical prote
35	37	62.7	3119	2	T18414 protein 9377 - mal
36	37	62.7	5138	2	B96695 hypothetical prote
37	36	61.0	182	2	T16423 hypothetical prote
38	36	61.0	206	2	B48441 antigen (C-termina
39	36	61.0	239	1	BVEC tonB protein - Esc
40	36	61.0	239	2	G85705 hypothetical prote
41	36	61.0	240	2	S13257 tonB protein - Sal
42	36	61.0	243	2	T45505 membrane protein
43	36	61.0	255	2	S30280 tonB protein - Yer
44	36	61.0	265	2	T00765 hypothetical prote
45	36	61.0	266	2	T44781 tonB protein (impo

ALIGNMENTS

RESULT 1
S35331
apidaecin 22 precursor - honeybee
C:Species: Apis mellifera (honeybee)
C>Date: 03-Feb-1994 #sequence_revision 03-Feb-1994 #text_change 21-Jul-2000
C:Accession: S35331
R:Castels-Josson, K.; Capaci, T.; Castels, P.; Tempst, P.
EMBO J. 12, 1569-1578, 1993
A>Title: Apidaecin multipeptide precursor structure: a putative mechanism for amplifi
A:Reference number: S35330; MUID:93223697
A:Accession: S35331
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-144 <CAS>
A:Cross-references: EMBL:X72576; NID:9297064; PIDN:CAAS1168.1; PID:9297065
C:Superfamily: procyclic acidic repetitive protein

Query Match 71.2%; Score 42; DB 2; Length 144;
Best Local Similarity 100.0%; Pred. No. 2.2;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 EAPEAPE 9
DB 35 EAPEAPE 42

RESULT 2
S35332
apidaecin 73 precursor - honeybee (fragment)
N:Contains: apidaecin 1a
C:Species: Apis mellifera (honeybee)
C>Date: 03-Feb-1994 #sequence_revision 03-Feb-1994 #text_change 03-Nov-2000
C:Accession: S35332; S05383
R:Castels-Josson, K.; Capaci, T.; Castels, P.; Tempst, P.
EMBO J. 12, 1569-1578, 1993
A>Title: Apidaecin multipeptide precursor structure: a putative mechanism for amplifi
A:Reference number: S35330; MUID:93223697
A:Accession: S35332
A:Molecule type: mRNA
A:Residues: 1-283 <CAS>
A:Cross-references: EMBL:X72577; NID:9297066; PIDN:CAAS1169.1; PID:94539289
A:Accession: S05383
A:Molecule type: protein
A:Residues: 258-283 <CAS>
C:Superfamily: proline-rich protein
F:266-283/Product: apidaecin 1a #status experimental <MAT>

Query Match 71.2%; Score 42; DB 2; Length 283;
Best Local Similarity 100.0%; Pred. No. 4.5;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 EAEPEAP 9
 |||||
 Db 62 EAEPEAP 69

RESULT 3
 S34224

Cyclin B - long-tailed hamster
 C:Species: Cricetus longicaudatus (long-tailed hamster)
 C:Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 16-Jul-1999
 C:Accession: S34224
 R:Maxiewicz, D.A.; Flick, M.B.; Mushel, R.J.; McKenna, W.G.
 submitted to the EMBL Data Library, March 1992
 A:Description: New features of mammalian cyclins seen in rat and chinese hamster cyclin
 A:Reference number: S34224
 A:Accession: S34224
 A:Status: preliminary
 A:Molecule type: mRNA
 A:Residues: 1-429 <MAR>
 A:Cross-references: EMBL:X64588; NID:g313764; PIDN:CAA45876.1; PID:g313765
 C:Superfamily: cyclin
 C:Keywords: cell cycle control

Query Match 67.8%; Score 40; DB 2; Length 429;
 Best Local Similarity 70.0%; Pred. No. 16;
 Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

OY 2 EAEPEAP 11
 |||||
 Db 98 EPEPEPEPV 107

RESULT 4

hypothetical protein SPBC11G11.05 - fission yeast (Schizosaccharomyces pombe)
 C:Species: Schizosaccharomyces pombe
 C:Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 03-Dec-1999
 C:Accession: T39332
 R:Saunders, D.; Harris, D.; McDougall, R.C.; Rajandream, M.A.; Barrell, B.G.
 submitted to the EMBL Data Library, October 1999
 A:Reference number: T39332
 A:Accession: T39332
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-251 <SAU>
 A:Cross-references: EMBL:ALJ37247; PIDN:CA859807.1; GSPDB:GN00067; SPDB:SPBC11G11.05
 A:Experimental source: strain 972h; cosmid c11G11
 C:Genetics:
 A:Gene: SPDB:SPBC11G11.05
 A:Map position: 2

Query Match 66.1%; Score 39; DB 2; Length 251;
 Best Local Similarity 87.5%; Pred. No. 13;
 Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 2 EAEPEAP 9
 |||||
 Db 185 ESEPEAP 192

RESULT 5

hematopoietic lineage cell-specific protein HSL - human
 C:Species: Homo sapiens (man)
 C:Date: 29-Jan-1993 #sequence_revision 29-Jan-1993 #text_change 05-Nov-1999
 C:Accession: S07633; A47478; B47478; D47478; E47478
 R:Klimura, D.; Kaneko, H.; Miyagoe, Y.; Akiyasu, T.; Watanabe, T.
 Nucleic Acids Res. 17, 9367-9379, 1989
 A:Title: Isolation and characterization of a novel human gene expressed specifically in
 A:Reference number: S07633; M0ID:90067934

A:Accession: S07633
 A:Molecule type: mRNA
 A:Residues: 1-486 <KIT>
 A:Cross-references: EMBL:X16663; NID:g32054; PIDN:CAA34651.1; PID:g32055
 R:Yamanashi, Y.; Okada, M.; Sema, T.; Yamori, T.; Umemori, H.; Tsunashima, S.; Toyosh
 Proc. Natl. Acad. Sci. U.S.A. 90, 3631-3635, 1993
 A:Title: Identification of HSL protein as a major substrate of protein-tyrosine kinases
 A:Reference number: A47478; M0ID:93234551
 A:Accession: A47478
 A:Status: preliminary

A:Molecule type: protein
 A:Residues: 4-19, 'XXX', '23-26', 'X', '79-93', 'X', '95', 'X', '134-146', '208-223', '274-284', 'X', '286', 'X'
 A:Experimental source: David, B-lymphoblastoid cells
 A>Note: sequence modified after extraction from NCBI backbone
 C:Superfamily: SH3 homology
 F:435-482/Domain: SH3 homology <SH3>

Query Match 66.1%; Score 39; DB 2; Length 486;
 Best Local Similarity 77.8%; Pred. No. 27;
 Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY 1 HEAEPEAP 9
 |||||
 Db 360 YEAEPEPEP 368

RESULT 6

hypothetical protein M03C11.2 - Caenorhabditis elegans
 C:Species: Caenorhabditis elegans
 C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999
 C:Accession: T23694
 R:McMurray, A.
 submitted to the EMBL Data Library, April 1995
 A:Reference number: T23694
 A:Accession: T23694
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-848 <MIT>
 A:Cross-references: EMBL:Z49128; PIDN:CA88959.1; GSPDB:GN00021; CESP:M03C11.2
 A:Experimental source: clone M03C11
 C:Genetics:
 A:Gene: CESP:M03C11.2
 A:Map position: 3
 A:introns: 113/2; 147/3; 185/3; 379/1; 482/3; 553/2; 688/3; 762/3

Query Match 66.1%; Score 39; DB 2; Length 848;
 Best Local Similarity 77.8%; Pred. No. 48;
 Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY 2 EAEPEAP 10
 |||||
 Db 506 EPEPEAP 514

RESULT 7

hypothetical protein F40F11.2 - Caenorhabditis elegans
 C:Species: Caenorhabditis elegans
 C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999
 C:Accession: T22028
 R:Dobson, R.
 submitted to the EMBL Data Library, May 1996
 A:Reference number: T22028
 A:Accession: T22028
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-1593 <MIT>
 A:Cross-references: EMBL:Z73426; NID:el343215; PIDN:CAA97793.1; GSPDB:GN00022; CESP:F
 A:Experimental source: clone F40F11
 C:Genetics:

A:Gene: CESP:F40P11.2
A:Map position: 4
A:Introns: 22/3; 96/3; 129/3; 206/1; 248/3; 710/1; 811/1; 1517/3; 1552/1

Query Match 66.1%; Score 39; DB 2; Length 1593;
Best Local Similarity 87.5%; Pred. No. 93;
Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 2 EAEPEAP 9
Db 851 ESEPEAP 858

RESULT 8
C72724
hypothetical protein APE0332 - Aeropyrum pernix (strain K1)
C:Species: Aeropyrum pernix
C>Date: 20-Aug-1999 #sequence_revision 20-Aug-1999 #text_change 09-Jun-2000
C:Accession: C72724
R:Kawarabayashi, Y.; Hino, Y.; Horikawa, H.; Yamazaki, S.; Halkawa, Y.; Jin-no, K.; Takahara, H.; Takamiya, M.; Masuda, S.; Funahashi, T.; Tanaka, T.; Kudoh, Y.; Yamazaki, J.; K DNA Res. 6; 83-101, 1999
A:Title: Complete genome sequence of an aerobic hyper-thermophilic Crenarchaeon, Aeropyrum pernix strain K1
A:Reference number: A72450; MUID:99310339
A:Accession: C72724
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-150 <KAW>
A:Cross-references: DDBJ:AP000059; NID:95103911; PIDN:BAA/9287.1; PID:Q1043073; PID:9510
A:Experimental source: strain K1
C:Genetics:
A:Gene: APE0332
C:Superfamily: Aeropyrum pernix hypothetical protein APE0332

Query Match 64.4%; Score 38; DB 2; Length 150;
Best Local Similarity 66.7%; Pred. No. 12;
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 3 AEPEAP 11
Db 45 AEPEAP 53

RESULT 9
S35330
apidaecin 14 precursor - honeybee
N:Contains: apidaecin II
C:Species: Apis mellifera (honeybee)
C>Date: 03-Feb-1994 #sequence_revision 03-Feb-1994 #text_change 21-Jul-2000
C:Accession: S35330; S06676
R:Castels-Josson, K.; Capaci, T.; Castels, P.; Tempst, P.
EMBO J. 12; 1569-1578, 1993
A:Title: Apidaecin multipeptide precursor structure: a putative mechanism for amplification of the honeybee defense system
A:Reference number: S35330; MUID:93223697
A:Accession: S35330
A:Molecule type: mRNA
A:Residues: 1-168 <CAS>
A:Cross-references: EMBL:X12575; NID:9297062; PIDN:CAA51167.1; PID:9297063
R:Castels, P.; Ampe, C.; Jacobs, F.; Vaack, M.; Tempst, P.
EMBO J. 8; 2387-2391, 1989
A:Title: Apidaecin: antibacterial peptides from honeybees.
A:Reference number: S05383; MUID:90005446
A:Accession: S06676
A:Molecule type: protein
A:Residues: 43-60 <CA2>
C:Superfamily: procytic acidic repetitive protein
F:43-60/Product: apidaecin II #status experimental <MAT>

Query Match 64.4%; Score 38; DB 2; Length 168;
Best Local Similarity 87.5%; Pred. No. 13;

Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 2 EAEPEAP 9
Db 35 EAEPEAP 42

RESULT 10
S33755
granzyme-like protein 1 (EC 3.4.21.-) precursor - rat
C:Species: Rattus norvegicus (Norway rat)
C>Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 22-Jun-1999
C:Accession: S33755
R:Amelrik, A.Y.; Varovoi, S.V.; Grigorenko, V.G.; Antonov, V.K.
FEBS Lett. 324; 226-230, 1993
A:Title: Identification, sequence analysis, and characterization of cDNA clones encoding rat granzyme-like protein 1
A:Reference number: S33755; MUID:93283341
A:Accession: S33755
A>Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-248 <AME>
A:Cross-references: EMBL:X66693; NID:9296177; PIDN:CAA47235.1; PID:9296178
C:Superfamily: trypsin; trypsin homology
C:Keywords: hydrolase; serine proteinase
F:21-241/Domain: trypsin homology <TRY>
F:65,109,204/Active site: His, Asp, Ser #status predicted

Query Match 64.4%; Score 38; DB 2; Length 248;
Best Local Similarity 54.5%; Pred. No. 20;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 1 HEAPEAP 11
Db 25 HEAPEAP 35

RESULT 11
PRH3
protease 3 (EC 3.4.21.-) precursor [validated] - human
N:Alternate names: AGP7; C-ANCA antigen; neutrophil proteinase 4; p29; Wegener's gran
N:Contains: myeloblastin
C:Species: Homo sapiens (man)
C>Date: 31-Mar-1993 #sequence_revision 31-Mar-1993 #text_change 08-Dec-2000
C:Accession: A45080; B46268; A43983; JH0331; A33751; S11091; A61176; A60148; A43982;
R:Sturrock, A.B.; Franklin, K.F.; Rao, G.; Marshall, B.C.; Rebentisch, M.B.; Lemons, J. Biol. Chem. 267; 21193-21199, 1992
A:Title: Structure, chromosomal assignment, and expression of the gene for proteinase 3
A:Reference number: A45080; MUID:93016043
A:Accession: A45080
A:Molecule type: DNA
A:Residues: 1-234; 'P' <STU>
A:Cross-references: GB:M97911
A:Note: sequence extracted from NCBI backbone (NCBIP:116205)
R:Zimmer, M.; Medcalf, R.L.; Fink, T.M.; Matmann, C.; Lichter, P.; Jenne, D.E.
Proc. Natl. Acad. Sci. U.S.A. 89; 8215-8219, 1992
A:Title: Three human elastase-like genes coordinately expressed in the myelomonocyte cell line U937
A:Reference number: A46268; MUID:92390417
A:Accession: B46268
A:Molecule type: DNA
A:Residues: 1-118; 'V', '120-134', 'AT', '137-256' <2IM>
A:Note: sequence extracted from NCBI backbone (NCBIN:112898, NCBIN:112900, NCBIN:1129
R:Labhaye, C.; Musette, P.; Cayre, Y.E.
Proc. Natl. Acad. Sci. U.S.A. 88; 9253-9256, 1991
A:Title: Wegener autoantigen and myeloblastin are encoded by a single mRNA.
A:Reference number: A43983; MUID:92021028
A:Accession: A43983
A:Molecule type: mRNA
A:Residues: 1-69; 'P', '71-256' <1A2>
A:Cross-references: GB:M75154; NID:9187398; PIDN:AAA5558.1; PID:9187399
R:Campanelli, D.; Weichior, M.; Fu, Y.; Nakata, M.; Shuman, H.; Nathan, C.; Gabay, J. J. Exp. Med. 172; 1709-1715, 1990
A:Title: Cloning of cDNA for proteinase 3: a serine protease, antibiotic, and autoant

A:Reference number: JH0331; MUID:91079774
 A:Accession: JH0331
 A:Molecule type: mRNA
 A:Residues: 'R',3-118, 'V',120-134, 'AT',137-254, 'P' <CAM>
 A:Cross-references: GB:X55668; NID:935687; PIDN:CAA39203.1; PID:9135280
 A:Note: part of this sequence, including the amino end of the mature protein, was confir
 R:Bories, D.; Raynal, M.C.; Solomon, D.H.; Darzynkiewicz, Z.; Cayre, Y.E.
 Cell 59, 959-968, 1989
 A:Title: Down-regulation of a serine protease, myeloblastin, causes growth arrest and d
 A:Reference number: A33751; MUID:90090622
 A:Accession: A33751
 A:Molecule type: mRNA
 A:Residues: 42-256 <BON>
 A:Cross-references: GB:M29142; NID:9188983; PIDN:AAA36342.1; PID:9188984
 A:Note: the authors translated the codon GGG for residue 49 as G10, GGC for residue 52 a
 R:Jenne, D.E.; Tschopp, J.; Luedemann, J.; Utecht, B.; Gross, W.L.
 Nature 346, 520, 1990
 A:Title: Wegener's antineutrophil proteinase.
 A:Reference number: S11091; MUID:90332035
 A:Accession: S11091
 A:Molecule type: mRNA
 A:Residues: 20-56 <JEN>
 R:Musette, P.; Labbaye, C.; Dorner, M.H.; Cayre, Y.E.; Casanova, J.L.; Kourilsky, P.
 Blood 77, 1398-1399, 1991
 A:Title: Wegener's antineutrophil proteinase and leukemia.
 A:Reference number: A61176; MUID:91159650
 A:Accession: A61176
 A:Molecule type: mRNA
 A:Residues: 1-42 <MUS>
 A:Cross-references: EMBL:X56606; NID:935189; PIDN:CAA39943.1; PID:935190
 APMIS 19(Suppl.), 26-27, 1990
 R:Goldschmidt, R.; Dolman, K.M.; Van Den Ende, M.E.; Van Der Meer-Gerritsen, C.H.; Son
 A:Title: The relation of 29 kD C-ANCA antigen to proteinase 3.
 A:Reference number: A60148; MUID:91136884
 A:Accession: A60148
 A:Molecule type: protein
 A:Residues: 28-48 <GOL>
 R:Rao, N.V.; Weber, N.G.; Marshall, B.C.; Gray, W.R.; Gray, B.H.; Hoidal, J.R.
 J. Biol. Chem. 266, 9540-9548, 1991
 A:Title: Characterization of proteinase-3 (PR-3), a neutrophil serine proteinase. Struct
 A:Reference number: A43982; MUID:91236723
 A:Accession: A43982
 A:Molecule type: protein
 A:Residues: 28-61, 'X',63, 'D',65-67,228-244 <RAO>
 R:Wilde, C.G.; Snable, J.L.; Griffith, J.E.; Scott, R.W.
 J. Biol. Chem. 265, 2038-2041, 1990
 A:Title: Characterization of two azurophil granule proteases with active-site homology t
 A:Reference number: A43981; MUID:90130450
 A:Accession: A43981
 A:Molecule type: protein
 A:Residues: 28-45, 'E',47,146-208, 'X',210-215, 'X',217-219 <WIL>
 R:Gibay, J.E.; Scott, R.W.; Campanelli, D.; Griffith, J.; Wilde, C.; Marra, M.N.; Seeger
 Proc. Natl. Acad. Sci. U.S.A. 86, 5610-5614, 1989
 A:Title: Antibiologic proteins of human polymorphonuclear leukocytes.
 A:Reference number: A33913; MUID:89315847
 A:Accession: C33913
 A:Molecule type: protein
 A:Residues: 28-45, 'E',47 <GAB>
 R:Niles, J.L.; McCluskey, R.T.; Almad, M.F.; Arnaout, M.A.
 Blood 74, 1888-1893, 1989
 A:Title: Wegener's granulomatosis antineutrophil proteinase is a novel neutrophil serine proteinase.
 A:Reference number: A60481; MUID:90028708
 A:Accession: A60481
 A:Molecule type: protein
 A:Residues: 28-38, 'X',40-47 <N12>
 R:Ohlsson, K.; Linder, C.; Rosengren, M.
 Biol. Chem. Hoppe-Seyler 371, 549-555, 1990
 A:Title: Monoclonal antibodies specific for neutrophil proteinase 4. Production and use
 A:Reference number: S10605; MUID:91025622
 A:Accession: S10605
 A:Molecule type: protein
 A:Residues: 28-52 <OHL>
 R:Luedemann, J.; Utecht, B.; Gross, W.L.

J. Exp. Med. 171, 357-362, 1990
 A:Title: Anti-neutrophil cytoplasm antibodies in Wegener's granulomatosis recognize a
 A:Reference number: PL0230; MUID:90111630
 A:Accession: PL0230
 A:Molecule type: protein
 A:Residues: 28-37, 'I',39-40, 'I',41-43 <LUE>
 C:Comment: This polymorphonuclear leukocyte serine protease from azurophilic granules
 C:Genetics:
 A:Gene: GDB:PRN3
 A:Cross-references: GDB:126876; OMIM:177020
 A:Map position: 19p13.3-19p13.3
 A:Introns: 21/1; 76/2; 123/3; 200/3
 C:Superfamily: trypsin; trypsin homology
 C:Keywords: glycoprotein; hydrolase; polymorphonuclear leukocyte; serine proteinase
 F:1-25/Domain: signal sequence #status predicted <SIG>
 F:26-27/Domain: propeptide #status predicted <PRO>
 F:28-256/Product: proteinase 3 #status experimental <MNT>
 F:28-243/Domain: trypsin homology <TRY>
 F:56-72,152-209,182-188,199-224/Disulfide bonds: #status predicted
 F:71,118,203/Active site: His, Asp, Ser #status predicted
 F:129,174/Binding site: carboxylate (Asn) (covalent) #status predicted

Query Match 64.4%; Score 38; DB 1; Length 256;
 Best Local Similarity 54.5%; Pred. No. 21;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAEPTM 11
 DB 32 HEAPHSRPFM 42

RESULT 12
 T51059
 hypothetical protein B12F1.110 [imported] - Neurospora crassa
 C:Species: Neurospora crassa
 C:Date: 21-Jul-2000 #sequence_revision 21-Jul-2000 #text_change 21-Jul-2000
 C:Accession: T51059
 R:Schlitz, U.; Aign, V.; Hohnsels, J.; Brandt, P.; Fartmann, B.; Holland, R.; Nyakatu
 submitted to the Protein Sequence Database, July 2000
 A:Reference number: 225286
 A:Accession: T51059
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-288 <SCH>
 A:Cross-references: EMBL:AL390091; GSPDB:GN00116; NCSP:B12F1.110
 C:Experimental source: BAC clone B12F1, strain OR74A
 C:Genetics:
 A:Gene: NCSP:B12F1.110
 A:Map position: 6
 A:Introns: 154/1

Query Match 64.4%; Score 38; DB 2; Length 288;
 Best Local Similarity 54.5%; Pred. No. 23;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAEPTM 11
 DB 58 HEHPESHPTL 68

RESULT 13
 S31848
 heat shock protein HSP30 - yeast (Saccharomyces cerevisiae)
 N:Alternate names: protein YCR021c
 C:Species: Saccharomyces cerevisiae
 C:Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 29-Oct-1999
 C:Accession: S31848; S19432; S30781
 R:Regnacq, M.; Boucherie, H.
 Curr. Genet. 23, 435-442, 1993
 A:Title: Isolation and sequence of HSP30, a yeast heat-shock gene coding for a hydrop
 A:Reference number: S31848; MUID:93306747

A:Accession: S31848
 A:Molecule type: DNA
 A:Residues: 1-263 <REG1>
 A:Cross-references: EMBL:M93123
 R:Feldmann, H.; Mannhaupt, G.; Vetter, I.
 submitted to the Protein Sequence Database, March 1992
 A:Reference number: S19429
 A:Accession: S19432
 A:Molecule type: DNA
 A:Residues: 1-190, 'A', 192-332 <FEU>
 A:Cross-references: EMBL:X59720; NID:q1907116; PIDN:CAA42313.1; PID:e264485; PID:q190716
 R:Regnacy, M.; Boucherie, H.
 submitted to the EMBL Data Library, January 1993
 A:Reference number: S30781
 A:Accession: S30781
 A:Molecule type: DNA
 A:Residues: 1-152, 'GY', 164-165, 167, 'A', 169, 'NSNRGL', 170-240, 'VFNQ', 270, 278-280, 'FNVFWI'
 A:Cross-references: EMBL:M93123
 A:Note: the difference at the carboxyl end is due to a frameshift error
 C:Genetics:
 A:Gene: SGD:HSP30
 A:Cross-references: SGD:S0000615; MIPS:YCR021C
 A:Map position: 3R
 C:Keywords: membrane protein

Query Match 64.4%; Score 38; DB 2; Length 332;
 Best Local Similarity 87.5%; Pred. No. 27;
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 HEAPEAE 8
 |||||
 Db 318 HEPEAE 325

RESULT 14
 S69371
 duodenase - bovine (fragment)
 C:Species: Bos primigenius taurus (cattle)
 C:Date: 24-Jul-1998 #sequence_revision 24-Jul-1998 #text_change 21-Nov-1998
 C:Accession: S69371
 R:Zamolodchikova, T.S.; Vorolyntseva, T.I.; Antonov, V.K.
 Eur. J. Biochem. 227, 866-872, 1995
 A:Title: Duodenase, a new serine protease of unusual specificity from bovine duodenal mu
 A:Reference number: S69371; MUID:95172075
 A:Accession: S69371
 A:Molecule type: protein
 A:Residues: 1-21 <ZAK>
 C:Superfamily: trypsin; trypsin homology

Query Match 62.7%; Score 37; DB 2; Length 21;
 Best Local Similarity 54.5%; Pred. No. 2.3;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;
 QY 1 HEAPEAEPIM 11
 |||:|:|
 Db 5 HEAKPSRPYM 15

RESULT 15
 T36257
 hypothetical protein SCE68.07c - Streptomyces coelicolor
 C:Species: Streptomyces coelicolor
 C:Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 03-Dec-1999
 C:Accession: T36257
 R:Murphy, L.; Harris, D.; James, K.D.; Parkhill, J.; Barrell, B.G.; Rajandream, M.A.
 submitted to the EMBL Data Library, June 1999
 A:Reference number: Z21576
 A:Accession: T36257
 A:Status: preliminary; translated from GB/EMBL/DDBJ
 A:Molecule type: DNA
 A:Residues: 1-125 <MUR>

A:Cross-references: EMBL:AL079345; PIDN:CA945343.1; GSPDB:GN00070; SCOEDB:SCE68.07c
 A:Experimental source: strain A3(2)
 C:Genetics:
 A:Gene: SCOEDB:SCE68.07c

Query Match 62.7%; Score 37; DB 2; Length 125;
 Best Local Similarity 87.5%; Pred. No. 15;
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 2 EAPEEAP 9
 |||||
 Db 5 EAPEEAP 12

Search completed: January 4, 2002, 08:41:31
 Job time: 169 sec

THIS PAGE BLANK (USPTO)


```

alignment_scores:
  quality: 42.00
  ratio: 4.667
  percent similarity: 100.000
  percent identity: 88.889

alignment_block:
  US-09-444-281-27 x US-09-103-840A-1

Align seg 1/1 to: US-09-103-840A-1 from: 1 to: 4411529
      3 AlAGUpProGUAAlAGUpProIleMet 11
      |||||
      260536 GCTGAACCGAGGACGAGCCGATC 260562

seq_name: /cgs2_6/prodata/2/lna/5B_Comp.seq:US-08-479-537A-1

seq_documentation_block:
  Sequence 1, Application US/08479537A
  Patent No. 5861381
  GENERAL INFORMATION:
  APPLICANT: CHAMRON, Pierre
  APPLICANT: KIENY, Marie-Paule
  APPLICANT: LAFRE, Richard
  APPLICANT: HAREUYEN, Mara
  TITLE OF INVENTION: PHARMACEUTICAL COMPOSITION FOR THE
  TREATMENT OR PREVENTION OF A MALIGNANT TUMOR
  NUMBER OF SEQUENCES: 5
  CORRESPONDENCE ADDRESS:
  ADDRESSEE: BURNS, DOANE, SNECKER & MATHIS, L.L.P.
  STREET: P.O. Box 1404
  CITY: Alexandria
  STATE: Virginia
  COUNTRY: United States
  ZIP: 22313-1404
  COMPUTER READABLE FORM:
  MEDIUM TYPE: Floppy disk
  OPERATING SYSTEM: PC-DOS/MS-DOS
  SOFTWARE: PatentIn Release #1.0, Version #1.30
  CURRENT APPLICATION DATA:
  APPLICATION NUMBER: US/08-479,537A
  FILING DATE: 07-JUN-1995
  CLASSIFICATION: 514
  PRIOR APPLICATION DATA:
  APPLICATION NUMBER: FR 90/13101
  FILING DATE: 23-OCT-1990
  PRIOR APPLICATION DATA:
  APPLICATION NUMBER: WO PCT/FR91/00835
  FILING DATE: 23-OCT-1991
  PRIOR APPLICATION DATA:
  APPLICATION NUMBER: US 08/039,320
  FILING DATE: 04-APR-1993
  PRIOR APPLICATION DATA:
  APPLICATION NUMBER: US 08/403,576
  FILING DATE: 14-MAR-1995
  ATTORNEY/AGENT INFORMATION:
  NAME: Teskin, Robin L.
  REGISTRATION NUMBER: 35,030
  REFERENCE/DOCKET NUMBER: 017753-025
  TELECOMMUNICATION INFORMATION:
  TELEPHONE: (703) 836-6620
  TELEFAX: (703) 836-2021
  INFORMATION FOR SEQ ID NO: 1:
  SEQUENCE CHARACTERISTICS:
  LENGTH: 6192 base pairs
  TYPE: nucleic acid
  STRANDEDNESS: single
  TOPOLOGY: linear
  MOLECULE TYPE: DNA (genomic)
  FEATURE:
  NAME/KEY: sig_peptide
  LOCATION: 58..120
  FEATURE:

```

```

1  NAME/KEY: repeat_region
2  LOCATION: 439..5239
3  OTHER INFORMATION: /note= "The nucleotides spanning
4  OTHER INFORMATION: 439-5239 constitute a repeated region wherein the repeat 1
5  OTHER INFORMATION: nucleotides and encodes 20 amino acids, 17 of which are fi
6  OTHER INFORMATION: The number of such repeats varies from 1 to 80."
7  FEATURE:
8  NAME/KEY: mat_peptide
9  LOCATION: 121..6166
10 FEATURE:
11 NAME/KEY: repeat_region
12 LOCATION: 457
13 OTHER INFORMATION: /note= "Nucleotide 457 is X1 = NNN
14 OTHER INFORMATION: which is the codon for Pro or Ala wherein Pro = CCT, CCC,
15 OTHER INFORMATION: or CCG; and Ala = GCT, GCC, GCA, or CCG."
16 FEATURE:
17 NAME/KEY: repeat_region
18 LOCATION: 487
19 OTHER INFORMATION: /note= "Nucleotide 487 is Y = NNN
20 OTHER INFORMATION: which is the codon for Thr or Asn wherein Thr = ACT, ACC,
21 OTHER INFORMATION: or ACG; and Asn = AAT or AAC."
22 FEATURE:
23 NAME/KEY: repeat_region
24 LOCATION: 496
25 OTHER INFORMATION: /note= "Nucleotide 496 is X2 = NNN
26 OTHER INFORMATION: which is the codon for Pro or Ala wherein Pro = CCT, CCC,
27 OTHER INFORMATION: or CCG; and Ala = GCT, GCC, GCA, or CCG."
28 US-08-479-537A-1
29
30 alignment_scores:
31     quality: 39.00      Length: 8
32     Ratio: 4.875      Gaps: 0
33     Percent similarity: 100.000      Percent identity: 87.500
34
35 alignment_block:
36 US-09-444-281-27 x US-08-479-537A-1/rev ..
37
38 Align seg 1/1 to reverse of: US-08-479-537A-1 from: 1 to: 6192
39
40 2 GUAAGAGUProGUAAGAGUPro 9
41  |||||
42 5306 GAAGCTGAGCCTGATGACAGAGCCT 5283
43
44 seq_name: /cgn2_6/ptodata/2/ina/6B_COMB.seq:US-09-083-116-1
45
46 seq_documentation_block:
47 : Sequence 1, Application US/09083116
48 : Patent No. 6203795
49 : GENERAL INFORMATION:
50 : APPLICANT: CHAMON, Pierre
51 : APPLICANT: KIENY, Marie-Paule
52 : APPLICANT: LAIHE, Richard
53 : APPLICANT: HAREUVENI, Mara
54 : TITLE OF INVENTION: PHARMACEUTICAL COMPOSITION FOR THE
55 : NUMBER OF SEQUENCES: 5
56 : CORRESPONDENCE ADDRESS:
57 : ADDRESSEE: BURNS, DOANE, SWECKER & MATHIS, L.L.P.
58 : STREET: P.O. Box 1404
59 : City: Alexandria
60 : STATE: Virginia
61 : COUNTRY: United States
62 : ZIP: 22313-1404
63
64 COMPUTER READABLE FORM:
65 MEDIUM TYPE: Floppy disk
66 COMPUTER: IBM PC compatible
67 OPERATING SYSTEM: PC-DOS/MS-DOS
68 SOFTWARE: Patentin Release #1.0, Version #1.30
69 CURRENT APPLICATION DATA:
70 APPLICATION NUMBER: US/09/083,116
71
72 FILING DATE:
73
74 CLASSIFICATION:

```



```

PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/479,537
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO PCT/FR91/00835
FILING DATE: 23-OCT-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/039,320
FILING DATE: 04-APR-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/403,576
FILING DATE: 14-MAR-1995
ATTORNEY/AGENT INFORMATION:
NAME: Teskin, Robin L.
REGISTRATION NUMBER: 35,030
REFERENCE/DOCKET NUMBER: 017753-025
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703) 836-6620
TELEFAX: (703) 836-2021
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 6192 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: sig_peptide
LOCATION: 58..120
FEATURE:
NAME/KEY: repeat_region
LOCATION: 439..5239
OTHER INFORMATION: /note= "The nucleotides spanning
OTHER INFORMATION: 439-5239 constitute a repeated region wherein the repeat is 6
OTHER INFORMATION: nucleotides and encodes 20 amino acids, 17 of which are fixed
OTHER INFORMATION: The number of such repeats varies from 1 to 80."
FEATURE:
NAME/KEY: mat_peptide
LOCATION: 121..6166
FEATURE:
NAME/KEY: repeat_region
LOCATION: 457
OTHER INFORMATION: /note= "Nucleotide 457 is X1 = NNN
OTHER INFORMATION: which is the codon for Pro or Ala wherein Pro = CCT, CCC, CCA
OTHER INFORMATION: or CCG; and Ala = GCT, GCC, GCA, or GCG."
FEATURE:
NAME/KEY: repeat_region
LOCATION: 487
OTHER INFORMATION: /note= "Nucleotide 487 is Y = NNN
OTHER INFORMATION: which is the codon for Thr or Asn wherein Thr = ACT, ACC, ACA
OTHER INFORMATION: or ACG; and Asn = AAT or AAC."
FEATURE:
NAME/KEY: repeat_region
LOCATION: 496
OTHER INFORMATION: /note= "Nucleotide 496 is X2 = NNN
OTHER INFORMATION: which is the codon for Pro or Ala wherein Pro = CCT, CCC, CCA
OTHER INFORMATION: or CCG; and Ala = GCT, GCC, GCA, or GCG."
US-09-083-116-1

alignment_scores:
Quality: 39.00 Length: 8
Ratio: 4.875 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 87.500

alignment_block:
US-09-444-281-27 x US-09-083-116-1/rev
Align seg 1/1 to reverse of: US-09-083-116-1 from: 1 to: 6192
2 GIUAlaGIUProGIUAlaGIUPro 9
|||||
5306 GAAGCTGAGCCTGATGACAGAGCCT 5283

```

```

seq_name: /cgn2_6/prodata/2/lna/58_COMB.seq:US-08-479-537A-4
seq_documentation_block:
; Sequence 4, Application US/08479537A
; Patent No. 5861381
; GENERAL INFORMATION:
; APPLICANT: CHAMBON, Pierre
; APPLICANT: KIENEY, Marie-Paule
; APPLICANT: LATHIE, Richard
; APPLICANT: HAREUVENI, Maira
; TITLE OF INVENTION: PHARMACEUTICAL COMPOSITION FOR THE
; TITLE OF INVENTION: TREATMENT OR PREVENTION OF A MALIGNANT TUMOR
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BURNS, DOANE, SWECKER & MATHIS, L.L.P.
; STREET: P.O. Box 1404
; City: Alexandria
; STATE: Virginia
; COUNTRY: United States
; ZIP: 22313-1404
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/479,537A
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: FR 90/13101
; FILING DATE: 23-OCT-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: WO PCT/FR91/00835
; FILING DATE: 23-OCT-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/039,320
; FILING DATE: 04-APR-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/403,576
; FILING DATE: 14-MAR-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Teskin, Robin L.
; REGISTRATION NUMBER: 35,030
; REFERENCE/DOCKET NUMBER: 017753-025
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703) 836-6620
; TELEFAX: (703) 836-2021
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 6449 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: sig_peptide
; LOCATION: 58..120
; FEATURE:
; NAME/KEY: repeat_region
; LOCATION: 439..5239
; OTHER INFORMATION: /note= "The nucleotides spanning
; OTHER INFORMATION: 439-5239 constitute a repeated region wherein the repeat 1
; OTHER INFORMATION: nucleotides and encodes 20 amino acids, 17 of which are fi
; OTHER INFORMATION: The number of such repeats varies from 1 to 80."
; FEATURE:
; NAME/KEY: mat_peptide
; LOCATION: 121..5661
; FEATURE:
; NAME/KEY: repeat_region
; LOCATION: 457
; OTHER INFORMATION: /note= "Nucleotide 457 is X1 = NNN

```

```

1  OTHER INFORMATION:  which is the codon for Pro or Ala wherein Pro = CCT, CCC, CCA
2  OTHER INFORMATION:  or CCG; and Ala = GCT, GCC, GCA, or GCG."
3  FEATURE:
4  NAME/KEY:  repeat_region
5  LOCATION:  487
6  OTHER INFORMATION:  /note= "Nucleotide 487 is Y = NNN
7  OTHER INFORMATION:  which is the codon for Thr or Asn wherein Thr = ACT, ACC, ACA
8  OTHER INFORMATION:  or ACG; and Asn = AAT or AAC."
9  FEATURE:
10 NAME/KEY:  repeat_region
11 LOCATION:  496
12 OTHER INFORMATION:  /note= "Nucleotide 496 is X2 = NNN
13 OTHER INFORMATION:  which is the codon for Pro or Ala wherein Pro = CCT, CCC, CCA
14 OTHER INFORMATION:  or CCG; and Ala = GCT, GCC, GCA, or GCG."
15 US-08-479-537A-4
16
17 alignment_scores:
18     quality: 39.00      length: 8
19     ratio: 4.875      gaps: 0
20     Percent Similarity: 100.000      Percent Identity: 87.500
21
22 alignment_block:
23 US-09-444-281-27 x US-08-479-537A-4/rev
24
25 Align seg 1/1 to reverse of: US-08-479-537A-4 from: 1 to: 6449
26
27     2  GUAlAGluProGUAlAGluPro 9
28     |||||||:::|||||||
29     5306 GAAGCTGAGCTGATGCAGAGCCT 5283
30
31 seq_name: /cgm2_6/prodata/2/ina/6B_COMB.seq:US-09-083-116-4
32
33 seq_documentation_block:
34 ; Sequence 4, Application US/09083116
35 ; Patent No. 6203795
36 ;
37 ; GENERAL INFORMATION:
38 ;
39 ; APPLICANT: CHAMON, Pierre
40 ; APPLICANT: KIENY, Marie-Paule
41 ; APPLICANT: LATHE, Richard
42 ; APPLICANT: HAREUVENI, Maira
43 ; TITLE OF INVENTION: PHARMACEUTICAL COMPOSITION FOR THE
44 ; TITLE OF INVENTION: TREATMENT OR PREVENTION OF A MALIGNANT TUMOR
45 ; NUMBER OF SEQUENCES: 5
46 ;
47 ; CORRESPONDENCE ADDRESS:
48 ; ADDRESSEE: BURNS, DOANE, SWECKER & MATHIS, L.L.P.
49 ; STREET: P.O. Box 1404
50 ; CITY: Alexandria
51 ; STATE: Virginia
52 ; COUNTRY: United States
53 ; ZIP: 22313-1404
54 ;
55 ; COMPUTER READABLE FORM:
56 ; MEDIUM TYPE: Floppy disk
57 ; COMPUTER: IBM PC compatible
58 ; OPERATING SYSTEM: PC-DOS/MS-DOS
59 ; SOFTWARE: Patent In Release #1.0, Version #1.30
60 ; CURRENT APPLICATION DATA:
61 ; APPLICATION NUMBER: US/09/083,116
62 ; FILING DATE:
63 ;
64 ; CLASSIFICATION:
65 ;
66 ; PRIOR APPLICATION DATA:
67 ; APPLICATION NUMBER: 08/479,537
68 ; FILING DATE:
69 ;
70 ; PRIOR APPLICATION DATA:
71 ; APPLICATION NUMBER: WO PCT/FR91/00835
72 ; FILING DATE: 23-OCT-1991
73 ;
74 ; PRIOR APPLICATION DATA:
75 ; APPLICATION NUMBER: US 08/039,320
76 ; FILING DATE: 04-APR-1993
77 ;
78 ; PRIOR APPLICATION DATA:
79 ; APPLICATION NUMBER: US 08/403,576
80 ; FILING DATE: 14-MAR-1995
81 ;
82 ATTORNEY/AGENT INFORMATION:

```

```

1 NAME: Teskin, Robin L.
2 REGISTRATION NUMBER: 35, 030
3 REFERENCE/DOCKET NUMBER: 017753-025
4 TELECOMMUNICATION INFORMATION:
5 TELEPHONE: (703) 836-6620
6 TELEFAX: (703) 836-2021
7 INFORMATION FOR SEQ ID NO: 4:
8 SEQUENCE CHARACTERISTICS:
9 LENGTH: 6449 base pairs
10 TYPE: nucleic acid
11 STRANDEDNESS: single
12 TOPOLOGY: linear
13 MOLECULE TYPE: DNA (genomic)
14 FEATURE:
15 NAME/KEY: sig_peptide
16 LOCATION: 58..120
17 FEATURE:
18 NAME/KEY: repeat_region
19 LOCATION: 439..5239
20 OTHER INFORMATION: /note="The nucleotides spanning
21 OTHER INFORMATION: 439-5239 constitute a repeated region wherein the repeat i
22 OTHER INFORMATION: nucleotides and encodes 20 amino acids, 17 of which are fi
23 OTHER INFORMATION: The number of such repeats varies from 1 to 80."
24 FEATURE:
25 NAME/KEY: mat_peptide
26 LOCATION: 121..5661
27 FEATURE:
28 NAME/KEY: repeat_region
29 LOCATION: 457
30 OTHER INFORMATION: /note="Nucleotide 457 is X1 = NNN
31 OTHER INFORMATION: which is the codon for Pro or Ala wherein Pro = CCT, CCC,
32 OTHER INFORMATION: or CCG; and Ala = GCT, GCC, GCA, or GCG."
33 FEATURE:
34 NAME/KEY: repeat_region
35 LOCATION: 487
36 OTHER INFORMATION: /note="Nucleotide 487 is Y = NNN
37 OTHER INFORMATION: which is the codon for Thr or Asn wherein Thr = ACT, ACC,
38 OTHER INFORMATION: or ACG; and Asn = AAT or AAC."
39 FEATURE:
40 NAME/KEY: repeat_region
41 LOCATION: 496
42 OTHER INFORMATION: /note="Nucleotide 496 is X2 = NNN
43 OTHER INFORMATION: which is the codon for Pro or Ala wherein Pro = CCT, CCC,
44 OTHER INFORMATION: or CCG; and Ala = GCT, GCC, GCA, or GCG."
45 US-09-083-116-4
46
47 alignment_scores:
48 Quality: 39.00 Length: 8
49 Ratio: 4.875 Gaps: 0
50 Percent Similarity: 100.000 Percent Identity: 87.500
51
52 alignment_block:
53 US-09-444-281-27 x US-09-083-116-4/rev ..
54
55 Align seg 1/1 to reverse of: US-09-083-116-4 from: 1 to: 6449
56
57 2 GtuatlagupProGtuatlagupPro 9
58 |||||||||:::|||||||
59 5306 GAAGCTGAGCCTGATGCAAGCCT 5283
60
61 seq_name: /cgn2_6/ptodata/2/ina/58_COMB.seq:us-08-394-600B-12
62
63 seq_documentation_block:
64 ; Sequence 12, Application US/08394600B
65 ; Patent No. 5845693
66 ; GENERAL INFORMATION:
67 ; APPLICANT: Halenbeck, Robert F.
68 ; APPLICANT: Jewell, David A.
69 ; APPLICANT: Koths, Kirston E.
70 ; APPLICANT: Kriegler, Michael
71 ; APPLICANT: Perez, Carl
72 TITLE OF INVENTION: Compositions for the inhibition of

```

TITLE OF INVENTION: Protein Hormone Formation and Uses Thereof
NUMBER OF SEQUENCES: 28
CORRESPONDENCE ADDRESS:
ADDRESSEE: McAndrews, Held & Malloy, Ltd.
STREET: 500 West Madison Street, 34th Floor
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60661
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/394,600B
FILING DATE: 02/27/95
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Donald J. Pochoplen
REGISTRATION NUMBER: 32,167
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312/707-8889
TELEFAX: 312/707-9155
TELEX:
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 83 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-394-600B-12

alignment_scores:
Quality: 38.00 Length: 11
Ratio: 3.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 54.545

alignment_block:

US-09-444-281-27 x US-08-394-600B-12/rev ..

Align seg 1/1 to reverse of: US-08-394-600B-12 from: 1 to: 83

1 HisGluAlaGluProGluAlaGluProIleMet 11
|||||
65 CACGAGCGCGCACACACTCCGCGCCCTACATG 33

seq_name: /cgn2_6/ptodata/2/1na/PCTUS_COMB.seq:PCT-US95-02513-12

seq_documentation_block:

Sequence 12, Application PC/TUS9502513

GENERAL INFORMATION:

APPLICANT: Halenbeck, Robert F.

APPLICANT: Jewell, David A.

APPLICANT: Koths, Kirston E.

APPLICANT: Krieglner, Michael

APPLICANT: Perez, Carl

TITLE OF INVENTION: Compositions for the Inhibition of

TITLE OF INVENTION: Protein Hormone Formation and Uses Thereof

NUMBER OF SEQUENCES: 22

CORRESPONDENCE ADDRESS:

ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &

ADDRESS: Borun

CITY: Chicago

STATE: Illinois

COUNTRY: United States of America

ZIP: 60606-6402

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/02513
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Williams Jr., Joseph A.
REGISTRATION NUMBER: 38,659
REFERENCE/DOCKET NUMBER: 27527/32404
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312/474-6300
TELEFAX: 312/474-0448
TELEX: 25-3856
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 83 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
PCT-US95-02513-12

alignment_scores:
Quality: 38.00 Length: 11
Ratio: 3.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 54.545

alignment_block:

US-09-444-281-27 x PCT-US95-02513-12/rev ..

Align seg 1/1 to reverse of: PCT-US95-02513-12 from: 1 to: 83

1 HisGluAlaGluProGluAlaGluProIleMet 11
|||||
65 CACGAGCGCGCACACACTCCGCGCCCTACATG 33

seq_name: /cgn2_6/ptodata/2/1na/5B_COMB.seq:US-08-394-600B-3

seq_documentation_block:

Sequence 3, Application US/08394600B

Patent No. 5843693

GENERAL INFORMATION:

APPLICANT: Halenbeck, Robert F.

APPLICANT: Jewell, David A.

APPLICANT: Koths, Kirston E.

APPLICANT: Krieglner, Michael

APPLICANT: Perez, Carl

TITLE OF INVENTION: Compositions for the Inhibition of

TITLE OF INVENTION: Protein Hormone Formation and Uses Thereof

NUMBER OF SEQUENCES: 28

CORRESPONDENCE ADDRESS:

ADDRESSEE: McAndrews, Held & Malloy, Ltd.

STREET: 500 West Madison Street, 34th Floor

CITY: Chicago

STATE: Illinois

COUNTRY: United States of America

ZIP: 60661

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/394,600B

FILING DATE: 02/27/95

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: Donald J. Pochoplen

REGISTRATION NUMBER: 32,167

REFERENCE/DOCKET NUMBER: 820,005/11850US05

TELECOMMUNICATION INFORMATION:
TELEPHONE: 312/707-8889
TELEFAX: 312/707-9155
TELEX:
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 771 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..771
US-08-394-600B-3

alignment_scores:
Quality: 38.00 Length: 11
Ratio: 3.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 54.545

alignment_block:

US-09-444-281-27 x US-08-394-600B-3 ..

Align seg 1/1 to: US-08-394-600B-3 from: 1 to: 771

1 HisGluAlaGluProGluAlaGluProIleMet 11
|||||||:|||||:|||||:|||||
94 CACGAGCGCAGCCACACCTCCGCCCTTACATG 126

seq_name: /cgn2_6/ptodata/2/ina/5B_COMB.seq:US-08-394-600B-22

seq_documentation_block:

Sequence 22, Application US/08394600B
Patent No. 5843693

GENERAL INFORMATION:

APPLICANT: Halenbeck, Robert F.

APPLICANT: Jewell, David A.

APPLICANT: Koths, Kirston E.

APPLICANT: Krieglner, Michael

APPLICANT: Perez, Carl

TITLE OF INVENTION: Compositions for the Inhibition of

NUMBER OF SEQUENCES: 28

CORRESPONDENCE ADDRESS:

ADDRESSEE: McAndrews, Heid & Malloy, Ltd.

STREET: 500 West Madison Street, 34th Floor

CITY: Chicago

STATE: Illinois

COUNTRY: United States of America

ZIP: 60661

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/394,600B

FILING DATE: 02/27/95

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: Donald J. Pochopien

REGISTRATION NUMBER: 32,167

REFERENCE/DOCKET NUMBER: 820,005/11850US05

TELECOMMUNICATION INFORMATION:

TELEPHONE: 312/707-8889

TELEFAX: 312/707-9155

TELEX:
INFORMATION FOR SEQ ID NO: 22:
SEQUENCE CHARACTERISTICS:
LENGTH: 771 base pairs
TYPE: nucleic acid

STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..771
US-08-394-600B-22

alignment_scores:
Quality: 38.00 Length: 11
Ratio: 3.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 54.545

alignment_block:

US-09-444-281-27 x US-08-394-600B-22 ..

Align seg 1/1 to: US-08-394-600B-22 from: 1 to: 771

1 HisGluAlaGluProGluAlaGluProIleMet 11
|||||||:|||||:|||||:|||||
94 CACGAGCGCAGCCACACCTCCGCCCTTACATG 126

seq_name: /cgn2_6/ptodata/2/ina/5B_COMB.seq:US-08-230-428B-3

seq_documentation_block:

Sequence 3, Application US/08230428B
Patent No. 5998378

GENERAL INFORMATION:

APPLICANT: Krieglner, Michael

APPLICANT: Halenbeck, Robert F.

APPLICANT: Perez, Carl

APPLICANT: Jewell, David A.

APPLICANT: Koths, Kirston E.

TITLE OF INVENTION: Compositions for the Inhibition of TNF

NUMBER OF SEQUENCES: 14

CORRESPONDENCE ADDRESS:

ADDRESSEE: CHIRON CORPORATION Intellectual Property - R440

STREET: 4560 Horton Street, P.O. Box 8097

CITY: Emeryville

STATE: California

COUNTRY: United States of America

ZIP: 94662-8097

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/230,428B

FILING DATE: 19-APR-1994

CLASSIFICATION: 514

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 07/905,546

FILING DATE: 25-JUN-1992

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 07/395,253

FILING DATE: 16-AUG-1989

ATTORNEY/AGENT INFORMATION:

NAME: Savereide, Paul B.

REGISTRATION NUMBER: 36,914

REFERENCE/DOCKET NUMBER: 0820,004

TELECOMMUNICATION INFORMATION:

TELEPHONE: (510) 601-2718

TELEFAX: (510) 655-3542

INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 771 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA

FEATURE:
NAME/KEY: CDS
LOCATION: 1..768
US-08-230-428B-3

alignment_scores:

Quality: 38.00 Length: 11
Ratio: 3.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 54.545

alignment_block:

US-09-444-281-27 x US-08-230-428B-3

Align seg 1/1 to: US-08-230-428B-3 from: 1 to: 771

1 HisGUAAGluProGUAAGluProIleMet 11
|||||
94 CACGAGGCGCACACCTCCGCCCTACATG 126

seq_name: /cgn2_6/prodata/2/ina/PCTUS_COMB.seq:PCT-US95-02513-3

seq_documentation_block:

Sequence 3, Application PC/TUS9502513
GENERAL INFORMATION:
APPLICANT: Halenbeck, Robert F.
APPLICANT: Jewell, David A.
APPLICANT: Kolts, Kirston E.
APPLICANT: Krieglner, Michael
APPLICANT: Perez, Carl
TITLE OF INVENTION: Compositions for the Inhibition of
TITLE OF INVENTION: Protein Hormone Formation and Uses Thereof
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &
ADDRESSEE: Borun
STREET: 6300 Sears Tower, 233 South Wacker Drive
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60606-6402
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/02513
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Williams Jr., Joseph A.
REGISTRATION NUMBER: 38,659
REFERENCE/DOCKET NUMBER: 27527/32404
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312/474-6300
TELEFAX: 312/474-0448
TELEX: 25-3856
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 771 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..771
PCT-US95-02513-3

alignment_scores:

Quality: 38.00 Length: 11

Ratio: 3.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 54.545

alignment_block:

US-09-444-281-27 x PCT-US95-02513-3

Align seg 1/1 to: PCT-US95-02513-3 from: 1 to: 771

1 HisGUAAGluProGUAAGluProIleMet 11
|||||
94 CACGAGGCGCACACCTCCGCCCTACATG 126

seq_name: /cgn2_6/prodata/2/ina/PCTUS_COMB.seq:PCT-US95-02513-22

seq_documentation_block:

Sequence 22, Application PC/TUS9502513
GENERAL INFORMATION:
APPLICANT: Halenbeck, Robert F.
APPLICANT: Jewell, David A.
APPLICANT: Kolts, Kirston E.
APPLICANT: Krieglner, Michael
APPLICANT: Perez, Carl
TITLE OF INVENTION: Compositions for the Inhibition of
TITLE OF INVENTION: Protein Hormone Formation and Uses Thereof
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &
ADDRESSEE: Borun
STREET: 6300 Sears Tower, 233 South Wacker Drive
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60606-6402
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/02513
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Williams Jr., Joseph A.
REGISTRATION NUMBER: 38,659
REFERENCE/DOCKET NUMBER: 27527/32404
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312/474-6300
TELEFAX: 312/474-0448
TELEX: 25-3856
INFORMATION FOR SEQ ID NO: 22:
SEQUENCE CHARACTERISTICS:
LENGTH: 771 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..771
PCT-US95-02513-22

alignment_scores:

Quality: 38.00 Length: 11
Ratio: 3.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 54.545

alignment_block:

US-09-444-281-27 x PCT-US95-02513-22

Align seg 1/1 to: PCT-US95-02513-22 from: 1 to: 771


```

: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 08/911,312
: FILING DATE: 14-AUG-1997
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 08/912,951
: FILING DATE: 14-AUG-1997
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 08/915,503
: FILING DATE: 14-AUG-1997
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: WO PCT/US97/17618
: FILING DATE: 01-OCT-1997
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: WO PCT/US97/17885
: FILING DATE: 01-OCT-1997
: ATTORNEY/AGENT INFORMATION:
: NAME: Apple, Randolph Ted
: REGISTRATION NUMBER: 36,429
: REFERENCE/DOCKET NUMBER: 015389-002610US
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (415) 576-0200
: TELEFAX: (415) 576-0300
: INFORMATION FOR SEQ ID NO: 646:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 90 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: DNA
: FEATURE:
: NAME/KEY: 1..90
: LOCATION: 1..90
: OTHER INFORMATION: /note="oligonucleotide 2T"
:
: US-08-974-549A-646

```

```

alignment_scores:
    Quality: 37.00      Length: 9
    Ratio: 4.111      Gaps: 0
    Percent Similarity: 100.000    Percent Identity: 66.667

```

alignment_block:
US-09-444-281-27 x US-08-974-549A-646/rev ..

Align seg 1/1 to reverse of: "US-08-974-549A-646 from: 1 to: 90

```

1 HisGluAlaGluProGluAlaGluPro 9
|||||
64 CACACGCTGAACCGACGCGAACCCCT 38

```

THIS PAGE BLANK (USPTO)

OM of: US-09-444-281-27 to: GenEmbl:* out_format: pfs
Date: Jan 4, 2002 10:54 AM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-2000 CompuGen Ltd.

Command line parameters:
-MODE=frame-p2n.model -DEV=slp
-O=/cgn2_1/USPRO.spool/US09444281/runat_04012002_084143_16195/app-query.fasta_1.210
-DB=GenEmbl -OPT=fastap -SUFFIX=trge -GAPOP=12.000 -GAPEXT=4.500
-MINMATCH=0.100 -LOOPEL=0.000 -LOOPEXT=0.000 -OGAPOP=4.500
-OGAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500 -OGAPEXT=6.000
-FEAPEXT=7.000 -YGAPEXT=10.000 -YGAPEXT=0.500 -DELOP=6.000
-DELEXT=7.000 -START=1 -MATRIX=blissum62 -TRANS=humand0.cdi
-LIST=45 -DOCALLIGN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0
-ALIGN=15 -MODE=LOCAL -OUTFM=pfis -NORM=ext -MINLEN=0
-MAXLEN=2000000000 -USER=US09444281@cgn1_15145 -NCPU=6
-ICPU=3 -LONGLOG -NO_XLPHY -WAIT -THREADS=1

Search information block:

Query: US-09-444-281-27
Query length: 11
Database: GenEmbl:*
Database sequences: 1472140
Database length: 34134837
Search time (sec): 1728.450000

Sequence	Strd Orig	Zscore	EScore	len	Documentation
gb.ph:AF109874	-	44.00	911.29	645.10	38347 AF109874 Bacteriophage Tuc200
gb.pr:AL391821	-	46.00	95.12	5.0e+03	158408 AL391821 Human DNA sequence
gb.hlg:AC021058	+	44.00	94.76	5.3e+03	169671 AC021058 Homo sapiens chrom
gb.to:RRPXPRT	+	43.00	126.67	88.61	966 Y09047 R. rattus mRNA for pxf pr
gb.ba:AF127222	-	43.00	120.21	202.80	2501 AF127222 Pseudomonas fluoresc
gb.hlg:AC015790	-	43.00	96.71	4.1e+03	79795 AC015790 Homo sapiens clone F
gb.pr:AF138709	-	43.00	95.95	4.9e+03	96874 AF138709 Human DNA sequence
gb.hlg:AC019290	-	43.00	91.31	8.3e+03	176712 AC019290 Homo sapiens clone
gb.hlg:AC016270	-	43.00	90.66	8.8e+03	189022 AC016270 Homo sapiens clone
gb.in:MAAP1D22	+	42.00	126.65	88.78	607 X72376 A. mellifera Apid22 mRNA
gb.pl:AF034945	+	42.00	124.86	111.67	790 AF034945 Zea mays glycine-rich
gb.to:MMY12380	+	42.00	123.87	126.90	915 Y12380 M. musculus ITGAV gene, F
gb.in:MAAP1D73	+	42.00	123.20	138.29	1010 X72577 Apis mellifera Apid73 m
gb.in:DDU2746	-	42.00	121.03	182.60	1390 U66367 Dictyostelium discoiden
gb.pr:HSSTPKERK	-	42.00	120.57	193.53	1486 U72746 Dictyostelium discoiden
gb.to:AB01015509	-	42.00	115.93	351.11	2946 X97630 H. sapiens mRNA for ser1
gb.hlg:AC017989	-	42.00	113.21	497.53	4397 AB012588 Mus musculus Munc18-1
gb.ba:AE006932	+	42.00	106.60	1.2e+03	11643 AC017989 Drosophila melanog
gb.pr:U73647	+	42.00	98.46	3.3e+03	38639 AE006932 Mycobacterium tuberc
gb.ba:MSG18D5	+	42.00	98.27	3.4e+03	39730 U73647 Human Chromosome 11 C
gb.ba:MSG140	+	42.00	98.09	3.5e+03	40806 AD90659 Mycobacterium tubercul
gb.in:CELR23E7	+	42.00	97.33	3.8e+03	44635 AF026205 Caenorhabditis eleg
gb.hlg:AC087785	-	42.00	94.99	5.1e+03	65477 AC087785 Homo sapiens chrom
gb.hlg:AC090264	-	42.00	94.71	5.3e+03	67203 AC090264 Homo sapiens chrom
gb.hlg:AC010702	-	42.00	94.58	5.4e+03	68457 AC010702 Drosophila melanog
gb.hlg:AC010571	-	42.00	91.67	7.9e+03	105160 AC010571 Drosophila melanog
gb.hlg:AC011254	-	42.00	91.01	8.6e+03	115884 AC011254 Drosophila melanog
gb.hlg:AC091284	-	42.00	90.89	8.7e+03	117993 AC091284 Mus musculus clone
gb.hlg:AL450338	+	42.00	89.19	1.1e+04	151608 AL450338 Homo sapiens chrom
gb.hlg:AP0033780	+	42.00	89.14	1.1e+04	152607 AP0033780 Homo sapiens chrom
gb.hlg:AC013253	+	42.00	88.94	1.1e+04	157108 AC013253 Homo sapiens chrom
gb.pr:AC092730	-	42.00	88.83	1.1e+04	157505 AC092730 Homo sapiens chrom
gb.pr:AC005242	-	42.00	88.81	1.1e+04	157505 AC005242 Homo sapiens chrom
gb.hlg:AL358034	-	42.00	88.76	1.1e+04	161388 AL358034 Homo sapiens chrom
gb.in:AC008228	-	42.00	88.70	1.2e+04	162951 AC008228 Drosophila melanog
gb.hlg:AC022490	-	42.00	88.37	1.2e+04	170892 AC022490 Homo sapiens chrom
gb.hlg:AC023920	-	42.00	88.32	1.2e+04	174852 AC023920 Homo sapiens chrom
gb.hlg:AC023920	+	42.00	87.37	1.3e+04	192453 AC023920 Homo sapiens clone
gb.in:AE003594	+	42.00	84.33	2.0e+04	305516 AE003594 Drosophila melanog
gb.in:AE003587	+	42.00	84.35	2.0e+04	309155 AE003587 Drosophila melanog

gb.ba:AP002996	-	42.00	83.52	2.2e+04	349619 AP002996 Mesorhizobium lo
gb.un:M27461	-	41.00	122.64	148.45	687 M27461 Figure 3. DNA sequenc
gb.pr:AF172450	-	41.00	116.59	322.63	1676 AF172450 Homo sapiens optoi
gb.pr:AF172452	-	41.00	114.26	433.06	2363 AF172452 Homo sapiens optoi
seq_name: gb_ph:AF109874					
seq_documentation block:					
LOCUS	AF109874	38347 bp	DNA	circular	PHG 15-MAR-2001
DEFINITION	Bacteriophage Tuc2009, complete genome.				
ACCESSION	AF109874	L26219	L31348	L31364	L31366
VERSION	AF109874.2	GI:13346831			
KEYWORDS					
SOURCE	Bacteriophage Tuc2009.				
ORGANISM	Bacteriophage Tuc2009.				
REFERENCE	1 (bases 24100 to 24323; 33392 to 34000; 36107 to 38347)				
AUTHORS	Arendt,E.K., Daly,C., Fitzgerald,G.F. and van de Guchte,M.				
TITLE	Molecular characterization of lactococcal bacteriophage Tuc2009 and identification and analysis of genes encoding lysis, a putative holin, and two structural proteins				
JOURNAL	Appl. Environ. Microbiol. 60 (6), 1875-1883 (1994)				
MEDLINE	94304164				
PUBMED	8031083				
REFERENCE	2 (bases 1921 to 2781)				
AUTHORS	van de Guchte,M., Daly,C., Fitzgerald,G.F. and Arendt,E.K.				
TITLE	Identification of the putative repressor-encoding gene ci of the temperate lactococcal bacteriophage Tuc2009				
JOURNAL	Gene 144 (1), 93-95 (1994)				
MEDLINE	94299176				
PUBMED	8026765				
REFERENCE	3 (bases 37811 to 38347; 1 to 1335)				
AUTHORS	van de Guchte,M., Daly,C., Fitzgerald,G.F. and Arendt,E.K.				
TITLE	Identification of int and attP on the genome of lactococcal bacteriophage Tuc2009 and their use for site-specific plasmid integration in the chromosome of Tuc2009-resistant Lactococcus lactis M01363				
JOURNAL	Appl. Environ. Microbiol. 60 (7), 2324-2329 (1994)				
MEDLINE	94356466				
PUBMED	8074513				
REFERENCE	4 (bases 1 to 38347)				
AUTHORS	van Sinderen,D., van de Guchte,M., Seegers,J.F.M.L. and Fitzgerald,G.F.				
TITLE	Molecular analysis of the temperate lactococcal phage Tuc2009				
JOURNAL	Unpublished				
REFERENCE	5 (bases 4602 to 11597)				
AUTHORS	McGirth,S., Seegers,J.F.M.L., Fitzgerald,G.F., van Sinderen,D. and van de Guchte,M.				
TITLE	Direct Submission				
JOURNAL	Submitted (27-NOV-1998) Microbiology, University College Cork, College Road, Cork, Ireland				
REFERENCE	6 (bases 1 to 38347)				
AUTHORS	van Sinderen,D., van de Guchte,M., Seegers,J.F.M.L. and Fitzgerald,G.F.				
TITLE	Direct Submission				
JOURNAL	Submitted (16-FEB-2000) Microbiology, University College Cork, College Road, Cork, Ireland				
REMARK	Sequence update by submitter				
COMMENT	On Mar 15, 2001 this sequence version replaced gi:508612 gi:496281 gi:496276 gi:496278 gi:509671 gi:5001694.				
FEATURES	Location/Qualifiers				
source	1..38347				
CDS	/organism="Bacteriophage Tuc2009"				
	/db_xref="taxon:35241"				
	complement(94..1218)				
	/note="int: ori1"				
	/codon_start=1				
	/transl_table=11				
	/product="integrase"				
	/protein_id="AA32608.1"				
	/db_xref="GI:508613"				
	/translation="MATYOKRCKTQWYISRTKQGLPRLTKGSGFSKSDQAQEAAMDE SKLKKGFIYDPIKQISEYFKDMELYYKKNAIDEMTYKVEQGLTKLTKTYMPVVLISE				

CDS
 ITASSYORALNKAETHAKASTGKGFHTRVASTIOPLIEEGRLOKDFTRAVVKGNGD
 KAEDKRFNDEKCOLVDYERPNRLNPVSSPMLPTIISITMSRASEAPGLVMDIDEN
 NNITKCRRTNNRNKNGKGGKKPKPDAGIRPIVISTEMLKRPFRQOQLFESLSTK
 PIIDPVCYHPYKRTITTSALQNTLDBALKLNLSTPLVHGLRHTHSAVLVHGVDM
 TVSKRLGHASVAITQOQYTHIIELEKDKDKLIELLML"
 complement(1339. .1860)
 /note="orf2"
 /codon_start=1
 /transl_table=11
 /product="unknown"
 /protein_id="AAK19844.1"
 /db_xref="GI:13346832"
 /translation="MKNGKTPKAKPIYKRWIWIIVVIVAVISALGGGKSGT
 STSTSSSKIKTAEPKKETATPISEPMKNDYVNSAADKYKGLKEFOGVSVT
 ANFKGTDTVTEAGNFTDNOFQDTAKAVVNDENMAQQLNSGQYTFQAAGDGYMMSDG
 WVYLDENNGVYK"
 complement(1921. .2781)
 /note="repressor; orf4"
 /codon_start=1
 /transl_table=11
 /product="c12009"
 /protein_id="AA21825.1"
 /db_xref="GI:509672"
 /translation="MYEIQNKVSGKIDYKRSFGLSQEELAKKIGVKTITSVEV
 GTRSPKPOLIKLSEFVAIDFEPQDSTMMNVSILSEINKISSOLEEPRKIVL
 NMTNOLDONONKESKSVIPINKIPDLPIYSIKLLENFIMPSNTEWYEDDMV
 DVAILGRIAGLELDAVENEDGRPVPAHFLSARDYKWLAWDGSMEKITYGAVL
 IEAVPVSQGTGAVLFQDDCATLKKVYHEIDCLKLVINSKEFKDQPRATQNPAAVI
 GGAVKVEIDL"
 3023. .3253
 /note="lambda cro analogue; orf5"
 /codon_start=1
 /transl_table=11
 /product="cro2009"
 /protein_id="AAK19845.1"
 /db_xref="GI:13346833"
 /translation="MAVEKELTALRADEKISKENAEILGTPETRYRKKEGSDMWG
 AMEFLASQNRIDIDFLDKKSTSGLEKAS"
 3267. .3983
 /note="mp3; orf6"
 /codon_start=1
 /transl_table=11
 /product="major structural protein"
 /protein_id="AAK19846.1"
 /db_xref="GI:13346834"
 /translation="MNOLITIQNNNDQVVSGBRLHFLGVTRYNDMPEDMVYGF
 TENVDFIGTEKRVPOGGRPSVDHALKLDMAKELISMIONNEKQKAOYFLEVYKEL
 KOQLPOTPEOQIALLARGNVNLKKVERIENSVDLTDREGIPNKAKVLOKAVASK
 VYMTFGKYSNAHKLLGAKVREFRYKDLNNRPDVYKSDIPLSRDEATEYLDMMQPS
 RPTLEIRGINSQTSFPEE"
 3996. .4331
 /note="orf7"
 /codon_start=1
 /transl_table=11
 /product="unknown"
 /protein_id="AAK19847.1"
 /db_xref="GI:13346835"
 /translation="MEQTLVQATISVLIPEDKILVDKVEYQELKEKDFDGVMDYV
 TKSNRISIPVYSKVLKRPDLRRKRSVENGGWVYVYVNGKDMWSPFRKEMDPINKFEY
 OFSGSGSGL"
 4328. .4456
 /note="orf8"
 /codon_start=1
 /transl_table=11
 /product="unknown"
 /protein_id="AAK19848.1"
 /db_xref="GI:13346836"
 /translation="MTYIYVNPETGETLFDLFDLITQNLRAKLAKKINAVLR"
 4469. .4663
 /note="orf9"
 /codon_start=1
 /transl_table=11
 /product="unknown"

CDS
 /protein_id="AAK19849.1"
 /db_xref="GI:13346837"
 /translation="MKEIEKLANNYKELIINKTSDLALKUNDGDIRKARKWLKQLFY
 TADRAETNELIKISTIDILDY"
 5038. .5253
 /note="orf11"
 /codon_start=1
 /transl_table=11
 /product="unknown"
 /protein_id="AAD37096.1"
 /db_xref="GI:5001696"
 /translation="MYTQIMNGREVLTPVYIGKYHNDLEKREVVGEVIESTYRRKDC
 TMYILRRSRTEREKAAMLNSCLSDMGY"
 5255. .5353
 /note="orf12"
 /codon_start=1
 /transl_table=11
 /product="unknown"
 /protein_id="AAD37097.1"
 /db_xref="GI:5001697"
 /translation="MSKQKSPAMHAEQVNSPKLLKLKLYHECLKI"
 5369. .5887
 /note="orf13"
 /codon_start=1
 /transl_table=11
 /product="unknown"
 /protein_id="AAD37098.1"
 /db_xref="GI:5001698"
 /translation="MTEKEPKYKYNKNDSELNVLGKYYKEHOJOCDEYEIOAESRKAI
 EEKYNALVEIEMERKLQAEQKEMDYFGLAHOYLLTLETSKPKRTINGSVPSFK
 EENYSDMLISELKEKGCGKITAKYTTESVDKAKAKAVIKQGGQLVSDGEIVGEF
 KEDKTEFFTVYV"
 5896. .6519
 /note="orf14"
 /codon_start=1
 /transl_table=11
 /product="unknown"
 /protein_id="AAD37099.1"
 /db_xref="GI:5001699"
 /translation="MPDKNSVRELNSIVSDKVEKKKSGKKNELSTLSYTMWATEK
 KKEPKATYEIKRKYSDGNELPYWDSTGTFWFTSVYVDVYHEMVLPMDAQANKAM
 KDKPKYKMYTNGEKSEVQASMEVDNKTIMRCLVKNIAFMGLIYIYAGEDLPEEPQ
 POLSDADLIEIKYKONPENKPNVDEFKTRSEHVAEMKAYIDWSK"
 6519. .6971
 /note="SSB; orf15"
 /codon_start=1
 /transl_table=11
 /product="single-strand binding protein"
 /protein_id="AAD37100.1"
 /db_xref="GI:5001700"
 /translation="MINNVYVGRIRPDPRLRHTPQNOAVGTGIAVNRQFRANKER
 EADLINCIVIRQQAENLAKPAKKGALIGTGRQTRNRYENQOGQYVYEVAVDTROM
 LESKRTQGOQTSFQAOONKKPQADDPKAPADPAGGTEISDDLPF"
 7096. .7878
 /note="Rep; orf16; contains origin of replication"
 /codon_start=1
 /transl_table=11
 /product="replication initiation protein"
 /protein_id="AAD37101.1"
 /db_xref="GI:5001701"
 /translation="MAQRRMEKSEVTTSDELVDMPSSQLLYFHLGMEADDEGFIGNA
 KMLSRAYGSNNDDLKLEAKGFVIAFPSSGVYVYKDWNLNNKIRKDRQKPTIYERKTL

alignment_scores: 46.00 Length: 9
 Ratio: 5.111 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:
 US-09-444-281-27 x AF109874/rev ..
 Align seg 1/1 to reverse of: AF109874 from: 1 to: 38347

```

seq_name: gb_pr:AL391821
seq_documentation_block:
LOCUS      AL391821    158408 bp        DNA             PRI          01-NOV-2000
DEFINITION Human DNA sequence from clone RP11-212B22 on chromosome X, complete
sequence.
ACCESSION  AL391821
VERSION    AL391821.7   GI:11121082
KEYWORDS   HTG.
SOURCE     human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 158408)
AUTHORS    Heath,P.
TITLE       Direct Submissions
JOURNAL     Submitted (31-OCT-2000) Sanger Centre, Hinxton, Cambridgeshire,
            CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk Clone
            requests: clonerequests@sanger.ac.uk
COMMENT     On Nov 8, 2000 this sequence version replaced gi:10944214.
            During sequence assembly data is compared from overlapping clones.
            Where difference assemblies are found these are annotated as variations
            together with a note of the overlapping clone name. Note that the
            variation annotation may not be found in the sequence submission
            corresponding to the overlapping clone, as we submit sequences with
            only a small overlap as described above.
            This sequence has been finished according to sequence map criteria
            as follows. An attempt is made to resolve all sequencing problems,
            such as compressions and repeats, but not necessarily within known
            annotated human repeat sequence elements (e.g. Alu). Where the
            sequence is ambiguous, there is an annotation using the 'unsure'
            feature key.
            The following abbreviations are used to associate primary accession
            numbers given in the feature table with their source databases:
            Em:, EMBL; Sw:, SWISSPROT; Tr:, TREMBL; Wp:, WORMEP; Information
            on the WormPEP database can be found at
            http://www.sanger.ac.uk/Projects/C_elegans/wormep/ This sequence
            was generated from part of bacterial clone contigs of human
            chromosome X, constructed by the Sanger Centre Chromosome X Mapping
            Group. Further information can be found at
            http://www.sanger.ac.uk/HGP/chrx
            RP11-212B22 is from the library RPCI-11.1 constructed at the
            Roswell Park Cancer Institute by the group of Pieter de Jong. For
            further details see http://bacpac.med.buffalo.edu/
            VECTOR: pBac3.6
            This sequence is the entire insert of clone RP11-212B22 The true
            left end of clone RP11-48719 is at 133439 in this sequence. The
            true right end of clone RP11-35GI8 is at 153997 in this sequence.
FEATURES             Location/Qualifiers
     source           1..158408
                     /organism="Homo sapiens"
                     /db_xref="taxon:9606"
                     /chromosome="X"
                     /clone="RP11-212B22"
                     /clone_id="RPCI-11.1"
     repeat_region    2..226
                     /note="LIM2 repeat: matches 5946..6163 of consensus"
     repeat_region    295..525
                     /note="WTNTH repeat: matches 229..494 of consensus"
     repeat_region    648..677
                     /note="15 copies 2 mer ca 90% conserved"
     repeat_region    2598..2649
                     /note="26 copies 2 mer ac 90% conserved"
     repeat_region    2780..2853
                     /note="37 copies 2 mer ct 74% conserved"
     misc_feature     complement(3446..3841)
                     /note="match: GSS: Em:A0136371"
                     4491..4655
                     /note="LIME repeat: matches 468..628 of consensus"

```

repeat_L_region	4804. .5008	/note="MP58C repeat: matches 3. .88 of consensus"
repeat_L_region	5044. .5181	/note="LM4 repeat: matches 1500. .1639 of consensus"
repeat_L_region	5271. .5414	/note="LM4 repeat: matches 2242. .2285 of consensus"
repeat_L_region	5420. .5718	/note="AluSq repeat: matches 1. .298 of consensus"
repeat_L_region	5726. .5994	/note="AluJo repeat: matches 1. .271 of consensus"
repeat_L_region	6011. .6787	/note="LM6C repeat: matches 2356. .2818 of consensus"
repeat_L_region	6801. .7262	/note="L1P4S repeat: matches 5682. .6143 of consensus"
repeat_L_region	7305. .7750	/note="LM1A9 repeat: matches 5860. .6308 of consensus"
repeat_L_region	7799. .8106	/note="LM4 repeat: matches 2841. .3151 of consensus"
repeat_L_region	8246. .8761	/note="LM4 repeat: matches 869. .1405 of consensus"
repeat_L_region	8819. .9030	/note="LM4 repeat: matches 1391. .1596 of consensus"
repeat_L_region	9073. .10246	/note="L1P23 repeat: matches 4979. .6146 of consensus"
repeat_L_region	10260. .10753	/note="LM4 repeat: matches 1575. .1608 of consensus"
repeat_L_region	11391. .11509	/note="FLM1C repeat: matches 1. .132 of consensus"
repeat_L_region	11510. .11821	/note="AluSq repeat: matches 1. .312 of consensus"
repeat_L_region	11834. .12407	/note="LM4 repeat: matches 2370. .2908 of consensus"
repeat_L_region	12408. .12684	/note="AluJo repeat: matches 1. .277 of consensus"
repeat_L_region	12685. .13112	/note="LM4 repeat: matches 2908. .3346 of consensus"
repeat_L_region	13191. .13493	/note="LM4 repeat: matches 3350. .3653 of consensus"
repeat_L_region	13515. .13538	/note="12 copies 2 mer at 100% conserved"
repeat_L_region	13539. .13894	/note="LM1A9 repeat: matches 4974. .5334 of consensus"
repeat_L_region	13855. .14205	/note="AluSq repeat: matches 1. .310 of consensus"
repeat_L_region	14206. .15193	/note="LM1A9 repeat: matches 5334. .6305 of consensus"
repeat_L_region	15269. .15357	/note="L2 repeat: matches 2644. .2748 of consensus"
repeat_L_region	15388. .15841	/note="L1P repeat: matches 4685. .5138 of consensus"
misc_feature	complement(15793. .16200)	/note="match: GSS: Em: A0034620"
repeat_L_region	16099. .16221	/note="L2 repeat: matches 2572. .2710 of consensus"
repeat_L_region	16260. .16732	/note="L2 repeat: matches 1992. .2403 of consensus"
repeat_L_region	17982. .18300	/note="AluSq repeat: matches 1. .307 of consensus"
repeat_L_region	18532. .18756	/note="MIR repeat: matches 14. .262 of consensus"
repeat_L_region	20534. .20587	/note="27 copies 2 mer tt 81% conserved"
repeat_L_region	21039. .21171	/note="AluSq/x repeat: matches 79. .209 of consensus"
repeat_L_region	22602. .22902	/note="AluSq repeat: matches 2. .302 of consensus"
repeat_L_region	23484. .23533	/note="25 copies 2 mer gt 96% conserved"
repeat_L_region	23532. .23561	/note="10 copies 3 mer gttg 90% conserved"
repeat_L_region	23877. .25040	/note="L2 repeat: matches 1141. .2323 of consensus"
repeat_L_region	25257. .25415	

```

repeat_region /note="L2 repeat: matches 2582. .2748 of consensus"
26553. .26594\
/note="21 copies: 2 mer aa 76% conserved"
misc_feature 26798. .27063
/note="match: GSS: Em: A0343061"
repeat_region 27948. .28106
/note="MER5B repeat: matches 10. .175 of consensus"
28430. .28618
/note="MER53 repeat: matches 1. .189 of consensus"
repeat_region 28941. .29026
/note="MIR repeat: matches 112. .198 of consensus"
30445. .30809
/note="THE1B repeat: matches 1. .364 of consensus"
repeat_region 31228. .31341
/note="L2 repeat: matches 2629. .2749 of consensus"
31996. .32079
/note="L2 repeat: matches 2620. .2698 of consensus"
repeat_region 32105. .32398
/note="AluY repeat: matches 1. .294 of consensus"
/note="complement(32313. .32867)"
misc_feature /note="match: GSS: Em: A0357777"
/note="complement(32386. .32876)"
repeat_region 32611. .32654
/note="match: GSS: Em: A0357649"
/note="22 copies 2 mer tt 79% conserved"
repeat_region 32804. .32877
/note="MER5A repeat: matches 21. .114 of consensus"
33673. .33723
/note="MIR repeat: matches 102. .152 of consensus"
/note="complement(34512. .34778)"
misc_feature /note="match: STS: Em: L24830"
35370. .35678
/note="AluY repeat: matches 1. .309 of consensus"
/note="complement(36574. .36908)"
misc_feature /note="match: STS: Em: HSC93A6"
36808. .37545
/note="L2 repeat: matches 2017. .2749 of consensus"
37906. .38033
/note="MER5A repeat: matches 59. .189 of consensus"
38390. .38732
/note="MIR repeat: matches 1. .365 of consensus"
38744. .38995
/note="MIR repeat: matches 8. .261 of consensus"
/note="complement(39579. .40134)"
misc_feature /note="match: STS: Em: HSC18H06"
39990. .40124
/note="L2 repeat: matches 2564. .2710 of consensus"
42729. .42884
/note="MER5A repeat: matches 4. .184 of consensus"
43123. .43367
/note="MIR repeat: matches 3. .261 of consensus"
repeat_region 44086. .44327
/note="AluSg/x repeat: matches 70. .308 of consensus"
45117. .45305
/note="MER53 repeat: matches 1. .194 of consensus"
45425. .45951
/note="L2 repeat: matches 1767. .2358 of consensus"
46025. .46336
/note="AluX repeat: matches 1. .312 of consensus"
46474. .46532
/note="MIR repeat: matches 65. .123 of consensus"

```

```

alignment_scores:
  Quality: 44.00      Length: 9
  Ratio: 5.500      Gaps: 0
  Percent Similarity: 88.889      Percent Identity: 88.889

```

```

alignment_block:
  US-09-444-281-27 x AL391821/rev

```

```

Align seg 1/1 to reverse of: AL391821 from: 1 to: 156408

```

```

1 HisC1uAgiuProGua1aGluPro 9

```

```

|||||
96328 CATGAGGCTGAGCCAGATTGAGCCT 96302
seq_name: gb_htg:AC021098

```

```

seq_documentation_block:
LOCUS AC021098 169671 bp DNA HTG 01-SEP-2000
DEFINITION Homo sapiens chromosome X clone RP11-33A2, WORKING DRAFT SEQUENCE,
18 unordered pieces.
ACCESSION AC021098.3 GI:7230836
VERSION AC021098
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 169671)
Waterston,R.H.
The sequence of Homo sapiens clone
2 (bases 1 to 169671)
Waterston,R.H.
Direct Submission
Submitted (14-JAN-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Mar 13, 2000 this sequence version replaced gi:6922906.

```

COMMENT

```

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H.NH0033A02
----- Summary Statistics -----

```

```

Sequencing vector: M13, 81%
Sequencing vector: Plasmid, 19%
Chemistry: Dye-primer ET; 81% of reads
Chemistry: Dye-terminator Big Dye; 19% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 159126 bases at least Q40
Consensus quality: 162659 bases at least Q30
Consensus quality: 164792 bases at least Q20
Insert size: 170000; agarose-fp
Insert size: 167971; sum-of-contigs
Quality coverage: 4.16 in Q20 bases; agarose-fp
Quality coverage: 4.21 in Q20 bases; sum-of-contigs

```

```

* NOTE: This is a 'working draft' sequence. It currently
* consists of 18 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

```

```

* 1 1196: contig of 1196 bp in length
* 1197 1296: gap of unknown length
* 1297 3284: contig of 1988 bp in length
* 3285 3384: gap of unknown length
* 3385 6023: contig of 2639 bp in length
* 6024 6123: gap of unknown length
* 6124 8768: contig of 2645 bp in length
* 8769 8868: gap of unknown length
* 8869 12171: contig of 3303 bp in length
* 12172 12271: gap of unknown length
* 12272 15977: contig of 3706 bp in length
* 15978 16077: gap of unknown length
* 16078 24250: contig of 8173 bp in length
* 24251 24350: gap of unknown length
* 24351 32681: contig of 8331 bp in length
* 32682 32781: gap of unknown length
* 32782 41939: contig of 9158 bp in length

```

```

* 41940 42039: gap of unknown length
* 42040 48802: contig of 6763 bp in length
* 48803 48902: gap of unknown length
* 48903 57427: contig of 8525 bp in length
* 57428 57527: gap of unknown length
* 57528 66323: contig of 8796 bp in length
* 66324 78283: gap of unknown length
* 78284 78384: gap of unknown length
* 78384 93662: contig of 15279 bp in length
* 93663 93762: gap of unknown length
* 93763 111526: contig of 17764 bp in length
* 111527 129714: contig of 18088 bp in length
* 129715 129815: gap of unknown length
* 129815 144537: contig of 14723 bp in length
* 144538 144638: gap of unknown length
* 144638 169671: contig of 25034 bp in length.
Location/Qualifiers
FEATURES
source
1. 169671
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="RP11-33A2"
1. 1196
/feature="assembly_name:Contig7"
1297. 3284
/feature="assembly_name:Contig8"
3385. 6023
/feature="assembly_name:Contig9"
6124. 8768
/feature="assembly_name:Contig10"
8869. 12171
/feature="assembly_name:Contig11"
12272. 15977
/feature="assembly_name:Contig12"
16078. 24250
/feature="assembly_name:Contig13"
24351. 32681
/feature="assembly_name:Contig14"
32782. 41939
/feature="assembly_name:Contig15"
42040. 48802
/feature="assembly_name:Contig16"
clone_end:Sp6
vector_side:right"
48903. 57427
/feature="assembly_name:Contig17"
clone_end:T7
vector_side:right"
57528. 66323
/feature="assembly_name:Contig18"
66424. 78283
/feature="assembly_name:Contig19"
78384. 93662
/feature="assembly_name:Contig20"
93763. 111526
/feature="assembly_name:Contig21"
111627. 129714
/feature="assembly_name:Contig22"
129815. 144537
/feature="assembly_name:Contig23"
144638. 169671
/feature="assembly_name:Contig24"

```

```

BASE COUNT 53919 a 31234 c 31915 g 50898 t 1705 others
ORIGIN

```

```

alignment_scores:
Quality: 44.00 Length: 9
Ratio: 5.500 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 88.889

```

```

alignment_block:
US-09-444-281-27 x AC021098

```

```

Align seg 1/1 to: AC021098 from: 1 to: 169671

```

```

1 HisGluaIaGluProGluAlaGluPro 9
|||||
44084 CATGACGCTGAGCCAGATTGACCT 44110

```

```

seq_name: gb_ro:RRXPFPRT

```

```

seq_documentation_block:
LOCUS RRPXPFPRT 966 bp mRNA ROD 01-OCT-1999
DEFINITION R. rattus mRNA for Pxf protein.
ACCESSION Y09049
VERSION Y09049.1 GI:6010290
KEYWORDS Pxf protein.
SOURCE black rat.
ORGANISM Rattus rattus
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
1 (bases 1 to 966)
AUTHORS Kammerer, S.
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 966)
AUTHORS Kammerer, S.
TITLE Direct Submission
SUBMITTED (22-OCT-1996) S. Kammerer, Kinderkrankenhaus, Labor f.
Molekulare Biologie, Lindwurmstr. 4, D-80337 Muenchen, FRG
Location/Qualifiers
FEATURES
source
1. 966
/organism="Rattus rattus"
/db_xref="taxon:10117"
/cell_type="myocardocyte"
<1. 75
/number=1
join(6. 75, 76. 185, 186. 351, 352. 437, 438. 599, 600. 776,
777. 821, 822. 905)
/feature="Pxf"
join(6. 75, 76. 185, 186. 351, 352. 437, 438. 599, 600. 776,
777. 821, 822. 905)
/feature="Pxf"
/codon_start=1
/product="Pxf protein"
/protein_id="CAA70258.1"
/db_xref="GI:6010291"
/translation="MAAEGCGAGVEADRELEELLESALDDFDRAKPSAPSPRTISA
PDASGPKRSQDITAKDALFASQEKFFQELFSELSAQTAEKAKMLAEPEHLV
EFOFKLSEAGVSGSDASSQOFTSLKFTLSGLAKNATDLDNGSSEELRKAREGL
GNDDEGEGENITLPIQMSLMONLSKDVLPSPAKETTERPPMLQSHQESIPPEQEKY
QOQHSYWGKICEQFEAETPTDEATHRAFEAVLDMQDLQDGHPPKELAGEMPGL
NFDLDLNLNSGPGANGEGCLIM"
76. 185
/feature="Pxf"
/number=2
186. 351
/feature="Pxf"
/number=3
352. 437
/feature="Pxf"
/number=4
438. 599
/feature="Pxf"
/number=5
600. 776
/feature="Pxf"
/number=6
777. 821
/feature="Pxf"
/number=7
822. 905
/feature="Pxf"
/number=8

```

```

exon
exon
exon
exon
exon

```


* Will be sequenced to completion. In the event that
* the record is updated, the accession number will
* be preserved.
1
* 902 1001: gap of 901 bp in length
* 1002 1010: contig of 909 bp in length
* 1911 2010: gap of 100 bp
* 2011 2938: contig of 928 bp in length
* 2939 3038: gap of 100 bp
* 3039 3033: contig of 895 bp in length
* 3934 4033: gap of 100 bp
* 4034 4931: contig of 898 bp in length
* 4932 5031: gap of 100 bp
* 5032 5905: contig of 874 bp in length
* 5906 6005: gap of 100 bp
* 6006 6884: contig of 889 bp in length
* 6895 6994: gap of 100 bp
* 6995 7902: contig of 908 bp in length
* 7903 8002: gap of 100 bp
* 8003 8887: contig of 885 bp in length
* 8888 9879: gap of 100 bp
* 9880 9979: gap of 100 bp
* 9980 10888: contig of 909 bp in length
* 10889 11908: contig of 920 bp in length
* 11909 12008: gap of 100 bp
* 12009 12888: contig of 880 bp in length
* 12889 12988: gap of 100 bp
* 12989 13882: contig of 894 bp in length
* 13883 13982: gap of 100 bp
* 13883 14889: contig of 907 bp in length
* 14890 14989: gap of 100 bp
* 14990 15897: contig of 908 bp in length
* 15898 15997: gap of 100 bp
* 15998 16890: contig of 893 bp in length
* 16891 16990: gap of 100 bp
* 16991 17891: contig of 901 bp in length
* 17892 17991: gap of 100 bp
* 17992 18891: contig of 900 bp in length
* 18892 18991: gap of 100 bp
* 18992 19894: contig of 903 bp in length
* 19895 19994: gap of 100 bp
* 19995 20860: contig of 866 bp in length
* 20861 20960: gap of 100 bp
* 20961 21867: contig of 907 bp in length
* 21868 21967: gap of 100 bp
* 21868 22868: contig of 901 bp in length
* 22869 22968: gap of 100 bp
* 22969 23859: contig of 891 bp in length
* 23860 23959: gap of 100 bp
* 23960 24832: contig of 873 bp in length
* 24833 24932: gap of 100 bp
* 24933 25833: contig of 901 bp in length
* 25834 25933: gap of 100 bp
* 25934 26845: contig of 912 bp in length
* 26846 26945: gap of 100 bp
* 26946 27854: contig of 909 bp in length
* 27855 27954: gap of 100 bp
* 27955 28864: contig of 910 bp in length
* 28865 28964: gap of 100 bp
* 28965 29856: contig of 892 bp in length
* 29857 29956: gap of 100 bp
* 29957 30835: contig of 879 bp in length
* 30836 30935: gap of 100 bp
* 30936 31843: contig of 908 bp in length
* 31844 31943: gap of 100 bp
* 31944 32864: contig of 921 bp in length
* 32865 32964: gap of 100 bp
* 32965 33849: contig of 885 bp in length
* 33850 33949: gap of 100 bp
* 33950 34823: contig of 874 bp in length
* 34824 34923: gap of 100 bp

* 34924 35815: contig of 892 bp in length
* 35816 35915: gap of 100 bp
* 35916 36826: contig of 911 bp in length
* 36827 36926: gap of 100 bp
* 36927 37836: contig of 910 bp in length
* 37837 37936: gap of 100 bp
* 37937 38837: contig of 901 bp in length
* 38838 38937: gap of 100 bp
* 38938 39818: contig of 881 bp in length
* 39819 39918: gap of 100 bp
* 39919 40829: contig of 911 bp in length
* 40830 40929: gap of 100 bp
* 40930 41819: contig of 890 bp in length
* 41820 41919: gap of 100 bp
* 41920 42789: contig of 870 bp in length
* 42790 42889: gap of 100 bp
* 42890 43767: contig of 878 bp in length
* 43768 43867: gap of 100 bp
* 43868 44753: contig of 886 bp in length
* 44754 44853: gap of 100 bp
* 44854 45777: contig of 924 bp in length
* 45778 45877: gap of 100 bp
* 45878 46780: contig of 903 bp in length
* 46781 46880: gap of 100 bp
* 46881 47802: contig of 922 bp in length
* 47803 47902: gap of 100 bp
* 47803 48803: contig of 901 bp in length
* 48804 48903: gap of 100 bp
* 48904 49807: contig of 904 bp in length
* 49808 49907: gap of 100 bp
* 49909 50790: contig of 883 bp in length
* 50791 50890: gap of 100 bp
* 50891 51810: contig of 920 bp in length
* 51811 51910: gap of 100 bp
* 51911 52812: contig of 902 bp in length
* 52813 52912: gap of 100 bp
* 52913 53796: contig of 884 bp in length
* 53797 53896: gap of 100 bp
* 53897 54818: contig of 922 bp in length
* 54819 54918: gap of 100 bp
* 54919 55819: contig of 901 bp in length
* 55820 55919: gap of 100 bp
* 55920 56836: contig of 917 bp in length
* 56837 56936: gap of 100 bp
* 56937 57850: contig of 914 bp in length
* 57851 57950: gap of 100 bp
* 57951 58874: contig of 924 bp in length
* 58875 58974: gap of 100 bp
* 58975 59847: contig of 873 bp in length
* 59848 59947: gap of 100 bp
* 59948 60838: contig of 891 bp in length
* 60839 60938: gap of 100 bp
* 60939 61829: contig of 891 bp in length
* 61830 61929: gap of 100 bp
* 61930 62806: contig of 877 bp in length
* 62807 62906: gap of 100 bp
* 62907 63784: contig of 878 bp in length
* 63785 63884: gap of 100 bp
* 63885 64792: contig of 908 bp in length
* 64793 64892: gap of 100 bp
* 64893 65789: contig of 897 bp in length
* 65790 65889: gap of 100 bp
* 65890 66771: contig of 882 bp in length
* 66772 66871: gap of 100 bp
* 66872 67778: contig of 907 bp in length
* 67779 67878: gap of 100 bp
* 67879 68780: contig of 902 bp in length
* 68781 68880: gap of 100 bp
* 68881 69764: contig of 884 bp in length
* 69765 69864: gap of 100 bp
* 69865 70743: contig of 879 bp in length
* 70744 70843: gap of 100 bp
* 70844 71756: contig of 913 bp in length

* 71757 71956: gap of 100 bp
* 71857 72761: contig of 905 bp in length

alignment_scores:
Quality: 43.00 Length: 10
Ratio: 4.300 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 80.000

alignment_block:

US-09-444-281-27 x AC015790/rev

Align seg 1/1 to reverse of: AC015790 from: 1 to: 79795

2 GlnA1agupProgluA1agupPro11exet 11

4914 GAAGCTCCGACGACGACGACCATCATG 4885

seq_name: gb.pr:AL138709

seq_documentation_block:

LOCUS AL138709 96874 bp DNA PRI 08-AUG-2001

DEFINITION Human DNA sequence from clone RP11-460G11 on chromosome 13,

complete sequence.

ACCESSION AL138709

VERSION AL138709.19 GI:15141998

KEYWORDS HTG.

SOURCE human.

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

TITLE 1 (bases 1 to 96874)

JOURNAL

COMMENT

On Aug 9, 2001 this sequence version replaced gi:14626037.

During sequence assembly data is compared from overlapping clones.

Where differences are found these are annotated as variations

together with a note of the overlapping clone name. Note that the

variation annotation may not be found in the sequence submission

corresponding to the overlapping clone, as we submit sequences with

only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all

regions were either double-stranded or sequenced with an alternate

chemistry or covered by high quality data (i.e., phred quality >=

30); an attempt was made to resolve all sequencing problems, such

as compressions and repeats: all regions were covered by at least

one plasmid subclone or more than one M13 subclone, and the

assembly was confirmed by restriction digest. The following

abbreviations are used to associate primary accession numbers given

in the feature table with their source databases: Em: EMBL; SW:

SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information on the WORMPEP

database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
Chromosome 13, constructed by the Sanger Centre Chromosome 13
Mapping Group. Further information can be found at
http://www.sanger.ac.uk/HGP/Chr13
RP11-460G11 is from the library RP11-11.2 constructed by the group
of Pieter de Jong. For further details see
http://www.chori.org/bacpac/home.htm
VECTOR: pACE3.6
IMPORTANT: This sequence is not the entire insert of clone
RP11-460G11 it may be shorter because we sequence overlapping
sections only once, except for a 100 base overlap.
The true right end of clone RP11-460G11 is at 96874 in this
sequence. The true right end of clone RP11-335618 is at 2000 in
this sequence.

FEATURES

Source

Location/Qualifiers
1..96874
/organism="Homo sapiens"
/db_xref="taxon:9606"

/chromosome="13"
/clone_id="RP11-460G11"
/clone_idb="RP11-11.2"
638..775
/note="AlusB/q repeat: matches 159..296 of consensus"
2734..2953
/note="MIR repeat: matches 20..252 of consensus"
3412..3570
/note="Char11c4 repeat: matches 1740..1907 of consensus"
4223..4292
/note="MIR repeat: matches 133..205 of consensus"
4354..4475
/note="MER5B repeat: matches 1..129 of consensus"
4980..5129
/note="MER5A repeat: matches 13..164 of consensus"
5146..5284
/note="MIR repeat: matches 9..148 of consensus"
5314..5483
/note="Char11c4 repeat: matches 17..174 of consensus"
5484..5788
/note="AluY repeat: matches 5..309 of consensus"
5789..6029
/note="Char11c4 repeat: matches 174..457 of consensus"
6536..6602
/note="L2 repeat: matches 2675..2743 of consensus"
6719..7125
/note="MLR2FB repeat: matches 4..414 of consensus"
7364..7568
/note="MIR repeat: matches 37..251 of consensus"
10015..10064
/note="MIR repeat: matches 41..90 of consensus"
10027..10085
/note="MER3 repeat: matches 144..204 of consensus"
10086..10325
/note="L1M1 repeat: matches 5550..5792 of consensus"
10326..10634
/note="AlusX repeat: matches 1..309 of consensus"
10635..11160
/note="L1M1 repeat: matches 5792..6304 of consensus"
11161..11179
/note="MER3 repeat: matches 130..144 of consensus"
11182..11347
/note="MER3 repeat: matches 1..167 of consensus"
11713..11780
/note="4 copies 17 mer 86% conserved"
11787..11881
/note="L2 repeat: matches 2575..2699 of consensus"
12183..12292
/note="MIR repeat: matches 126..246 of consensus"
13008..13053
/note="23 copies 2 mer tc 73% conserved"
13063..13218
/note="6 copies 26 mer 67% conserved"
13065..13178
/note="57 copies 2 mer tc 67% conserved"
13674..13759
/note="MIR repeat: matches 78..165 of consensus"
15493..15544
/note="26 copies 2 mer ac 90% conserved"
16508..17150
/note="MER41B repeat: matches 3..635 of consensus"
18228..18286
/note="tRNA-Ile-ATP repeat: matches 14..74 of consensus"
18765..19215
/note="L2 repeat: matches 2290..2748 of consensus"
19216..19985
/note="LTR39 repeat: matches 1..794 of consensus"
19986..20070
/note="L2 repeat: matches 2208..2290 of consensus"
21039..21339
/note="AlusG repeat: matches 1..301 of consensus"
21350..21658
/note="AlusG repeat: matches 1..307 of consensus"


```

repeat_region 21674..21731
/note="MER53 repeat: matches 127. .186 of consensus"
repeat_region 21732..22043
/note="AluSx repeat: matches 1. .312 of consensus"
repeat_region 22044..22175
/note="MER53 repeat: matches 6. .127 of consensus"
repeat_region 22862..23024
/note="MIR repeat: matches 87. .248 of consensus"
repeat_region 24844..25143
/note="AluSx repeat: matches 1. .302 of consensus"
repeat_region 26057..26200
/note="MER5A repeat: matches 17. .187 of consensus"
repeat_region 27843..28278
/note="MLTID repeat: matches 1. .466 of consensus"
repeat_region 28375..28436
/note="L2 repeat: matches 2645. .2710 of consensus"
repeat_region 30397..30480
/note="2 copies 42 mer 90% conserved"
repeat_region 31856..32054
/note="MIR repeat: matches 37. .261 of consensus"
repeat_region 32358..32528
/note="L2 repeat: matches 2092. .2279 of consensus"
repeat_region 32608..32648
/note="MLTID repeat: matches 201. .240 of consensus"
repeat_region 32649..32947
/note="AluJo repeat: matches 1. .294 of consensus"
repeat_region 32948..33144
/note="MLTID repeat: matches 1. .201 of consensus"
repeat_region 33183..33230
/note="L2 repeat: matches 2677. .2724 of consensus"
repeat_region 34446..34647
/note="MER2 repeat: matches 1. .211 of consensus"
repeat_region 34678..34757
/note="MER2 repeat: matches 264. .344 of consensus"
repeat_region 38187..38646
/note="MLTIC repeat: matches 6. .466 of consensus"
repeat_region 38848..38958
/note="MER81 repeat: matches 3. .114 of consensus"
repeat_region 40042..40152
/note="L2 repeat: matches 2381. .2492 of consensus"
repeat_region 41116..41251
/note="MIR repeat: matches 15. .151 of consensus"
repeat_region 42941..43367
/note="LTP168 repeat: matches 39. .464 of consensus"
repeat_region 44219..44302
/note="MLTID repeat: matches 242. .318 of consensus"
repeat_region 46089..47392
/note="LTPA2 repeat: matches 3642. .4945 of consensus"
repeat_region 47393..48587
/note="LTPA2 repeat: matches 4952. .6146 of consensus"
repeat_region 48647..48781
/note="MIR repeat: matches 85. .234 of consensus"
repeat_region 49096..49396
/note="AluSg repeat: matches 1. .296 of consensus"
repeat_region 49623..51400
/note="LTPA7 repeat: matches 4358. .6143 of consensus"
repeat_region 52702..53151
/note="MER39 repeat: matches 13. .455 of consensus"
repeat_region 53233..53451
/note="MER72 repeat: matches 445. .668 of consensus"
repeat_region 53452..53750
/note="AluSg repeat: matches 1. .294 of consensus"
repeat_region 53751..54215
/note="MER72 repeat: matches 1. .445 of consensus"
repeat_region 54223..54364
/note="L2 repeat: matches 2563. .2750 of consensus"
repeat_region 54866..55171
/note="AluY repeat: matches 2. .303 of consensus"
repeat_region 55239..55292
/note="27 copies 2 mer tt 70% conserved"
repeat_region 57830..58112
/note="AluSg repeat: matches 3. .304 of consensus"
repeat_region 58224..58516

```

```

alignment_scores:
  Quality: 43.00      Length: 11
  Ratio: 4.778      Gaps: 0
  Percent Similarity: 81.818      Percent Identity: 63.636

alignment_block:
US-09-444-281-27 x AL138709/rev ..

Align seg 1/1 to reverse of: AL138709 from: 1 to: 96874

1 HtsguAlagluProgluAlagluProleleMet 11
||||| ||||||| |||||||:
66249 CACGAGCAGACCTGACCCGAGCCCTGGTT 66217

seq_name: gb_hlg:AC019290

seq_documentation_block:
LOCUS AC019290 176712 bp DNA HTG 09-SEP-2000
DEFINITION Homo sapiens clone RP11-649M10, WORKING DRAFT SEQUENCE, 15
unordered pieces.
ACCESSION AC019290
VERSION AC019290.4 GI:10045469
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 176712)
REFERENCE Birren,B., Linton,L., Nusbaum,C. and Lander,E.
Homo sapiens, clone RP11-649M10
unpublished
2 (bases 1 to 176712)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
Anderson,S., Baldwin,J., Barna,N., Beckerly,R., Beda,F.,
Boguslavsky,L., Bouckhalter,B., Brown,A., Burkett,G., Castle,A.,
Choepel,Y., Colangelo,M., Collins,S., Collymore,A., Cooke,P.,
DeArlelano,K., Dewar,K., Domino,M., Doyle,M., Feneator,J.,
Perreira,P., Fitzhugh,W., Forrest,C., Gage,D., Galagan,J.,
Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,
Landers,T., Lehoczeky,J., Levine,R., Lieu,C., Liu,G., Locke,K.,
Macdonald,P., Margulis,N., McEwan,P., McGuirk,A., McKernan,K.,
McPheeters,R., Meldrum,J., Meneus,L., Morrow,J., Naylor,J.,
Norman,C.H., O'Connor,T., O'Donnell,P., Oliver,T.M., Peterson,K.,
Pierre,N., Pisanl,C., Pollara,V., Raymond,C., Riley,R., Rothman,D.,
Roy,A., Santos,R., Severy,P., Spencer,B., Stange-Thomann,N.,
Stojanovic,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
Tirrell,A., Vassiliev,H., Viel,R., Vo,A., Wu,X., Wyman,D., Ye,W.J.,
Zimmer,A. and Zody,M.
Direct Submission
Submitted (31-DEC-1999) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA.
On Sep 9, 2000 this sequence version replaced gi:6721321.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: MIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L4312
----- Summary Statistics
Sequencing vector: M13: M7815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 167161 bases at least Q40
Consensus quality: 171786 bases at least Q30
Consensus quality: 173554 bases at least Q20

```

Insert size: 182000; agarose-fp
 Insert size: 175312; sum-of-ctnigs
 Quality coverage: 4.3 in Q20 bases; agarose-fp
 Quality coverage: 4.5 in Q20 bases; sum-of-ctnigs

NOTE: This is a 'working draft' sequence. It currently consists of 15 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

```

1 31590: contig of 31590 bp in length
* 31591 31690: gap of 100 bp
* 31691 33087: contig of 1397 bp in length
* 33088 33187: gap of 100 bp
* 33188 34887: contig of 1700 bp in length
* 34888 34987: gap of 100 bp
* 34988 37644: contig of 2657 bp in length
* 37645 37744: gap of 100 bp
* 37745 40979: contig of 3235 bp in length
* 40980 41079: gap of 100 bp
* 41080 46341: contig of 5262 bp in length
* 46342 46441: gap of 100 bp
* 46442 53992: contig of 7551 bp in length
* 53993 54092: gap of 100 bp
* 54093 60115: contig of 6024 bp in length
* 60117 60216: gap of 100 bp
* 60217 68183: contig of 7967 bp in length
* 68184 68283: gap of 100 bp
* 68284 76928: contig of 8645 bp in length
* 76929 77028: gap of 100 bp
* 77029 95627: contig of 18599 bp in length
* 95628 95727: gap of 100 bp
* 95728 115794: contig of 20067 bp in length
* 115795 115894: gap of 100 bp
* 115895 13115: contig of 27221 bp in length
* 143116 143215: gap of 100 bp
* 143216 174379: contig of 31164 bp in length
* 174380 174479: gap of 100 bp
* 174480 176712: contig of 2233 bp in length.
  
```

FEATURES

```

Source
1. 176712
  /organism="Homo sapiens"
  /db_xref="taxon:9606"
  /clone="RP11-649M10"
  /clone_lib="RP11 Human Male BAC"
  1. 31590
    /note="assembly-fragment"
    clone_end:SP6
    vector_side:left"
  misc_feature
    31691..33087
    /note="assembly-fragment"
  misc_feature
    33188..34887
    /note="assembly-fragment"
  misc_feature
    34988..37644
    /note="assembly-fragment"
  misc_feature
    37745..40979
    /note="assembly-fragment"
  misc_feature
    41080..46341
    /note="assembly-fragment"
  misc_feature
    46442..53992
    /note="assembly-fragment"
  misc_feature
    54093..60116
    /note="assembly-fragment"
  misc_feature
    60217..68183
    /note="assembly-fragment"
  misc_feature
    68284..76928
    /note="assembly-fragment"
  misc_feature
    77029..95627
    /note="assembly-fragment"
  misc_feature
    95728..115794
  
```

```

misc_feature /note="assembly-fragment"
115895..143115
  /note="assembly-fragment"
misc_feature 143216..174379
  /note="assembly-fragment"
misc_feature 174480..176712
  /note="assembly-fragment"
  clone_end:17
  vector_side:right"
  
```

BASE COUNT 53093 a 35421 c 35789 g 51008 t 1401 others

alignment_scores:
 Quality: 43.00 Length: 11
 Ratio: 4.300 Gaps: 0
 Percent Similarity: 90.909 Percent Identity: 72.727

alignment_block:
 US-09-444-281-27 x AC019290/rev ..

Align seq 1/1 to reverse of: AC019290 from: 1 to: 176712

```

1 HisgluAlagluProgluAlagluProilemet 11
|||||:|||||:|||||:|||||
65329 CATGAGGCTCCGCCAGACAGCATGCCATCATG 65297
  
```

seq_name: gp_htg:AC016270

seq_documentation_block:

LOCUS AC016270 189022 bp DNA HTG 16-MAR-2000
 DEFINITION Homo sapiens clone RP11-17111, WORKING DRAFT SEQUENCE, 15 unordered pieces

AC016270
 AC016270.4 GI:7248975

VERSION HTG: HTGS_PHASE1; HTGS_DRAFT.

KEYWORDS human.

SOURCE

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

AUTHORS Birren,B., Linton,L., Nusbaum,C. and Lander,E.

TITLE Homo sapiens, clone RP11-17111

JOURNAL Unpublished

2 (bases 1 to 189022)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,

Baldwin,J., Barna,N., Beckert,K., Boguslavsky,L., Bouknight,B.,

Brown,A., Castle,A., Colangelo,M., Collins,S., Collymore,A.,

Cooke,P., DeArrellano,K., Dewar,K., Domino,M., Donelan,L., Doyle,M.,

Ferreira,P., FitzHugh,W., Forrest,C., Funke,R., Gage,D.,

Galagan,J., Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,

Howland,J.C., Johnson,R., Jones,C., Kann,L., Kartas,A., Klein,J.,

Lehocky,J., Liu,C., Locke,K., MacDonald,P., Marquis,N.,

McEwan,P., McGurk,A., McKernan,K., McLaughlin,J., Meldrum,J.,

Morrow,J., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,

Peterson,K., Pollara,V., Riley,R., Roy,A., Santos,R., Severi,P.,

Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,

Testaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X.,

Wymann,D., Ye,W.J., Zimmer,A. and Zody,M.

TITLE

JOURNAL

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

```

Center clone name: 17.1.11
----- Summary Statistics
Sequencing vector: M13; M77815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 182484 bases at least Q40
Consensus quality: 184881 bases at least Q40
Consensus quality: 186037 bases at least Q20
Insert size: 187620; agarose-1p
Insert size: 187622; sum-of-contigs
Quality coverage: 5.8 in Q20 bases; agarose-1p
Quality coverage: 5.8 in Q20 bases; sum-of-contigs
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 15 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
* 1 1585: contig of 1585 bp in length
* 1586 1685: gap of 100 bp
* 1586 3061: contig of 1376 bp in length
* 3062 3161: gap of 100 bp
* 3162 7504: contig of 4343 bp in length
* 7505 7604: gap of 100 bp
* 7605 10536: contig of 2932 bp in length
* 10537 10636: gap of 100 bp
* 10637 13824: contig of 3188 bp in length
* 13825 13924: gap of 100 bp
* 13925 18777: contig of 4853 bp in length
* 18778 18877: gap of 100 bp
* 18878 29087: contig of 10210 bp in length
* 29088 29187: gap of 100 bp
* 29188 40374: contig of 11187 bp in length
* 40375 40474: gap of 100 bp
* 40475 55299: contig of 14825 bp in length
* 55300 55399: gap of 100 bp
* 55400 67577: contig of 12178 bp in length
* 67578 67677: gap of 100 bp
* 67678 86116: contig of 18439 bp in length
* 86117 86216: gap of 100 bp
* 86217 104553: contig of 18337 bp in length
* 104554 104653: gap of 100 bp
* 104654 125313: contig of 20660 bp in length
* 125314 125413: gap of 100 bp
* 125414 154037: contig of 28624 bp in length
* 154038 154137: gap of 100 bp
* 154138 189022: contig of 34885 bp in length.
*
* Location/Qualifiers
*
* 1..189022
* /organism="Homo sapiens"
* /db_xref="taxon:9606"
* /clone_RP11-17111"
* /clone_11b="RPC1-11 Human Male BAC"
*
* 1..1585
* /note="assembly-fragment"
* 1686..3061
* /note="assembly-fragment"
* 3162..7504
* /note="assembly-fragment"
* 7605..10536
* /note="assembly-fragment"
* 10637..13824
* /note="assembly-fragment"
* 13925..18777
* /note="assembly-fragment"
* 18878..29087
* /note="assembly-fragment"
* 29188..40374
* /note="assembly-fragment"
* clone_end:5p6

```

```

misc_feature      vector_side:left
40475..55299
/note="assembly_fragment"
misc_feature      55400..67577
/note="assembly_fragment"
misc_feature      67678..86116
/note="assembly_fragment"
misc_feature      86217..104553
/note="assembly_fragment"
misc_feature      104654..125313
/note="assembly_fragment"
clone_end:"77
vector_side:right"
misc_feature      125414..154037
/note="assembly_fragment"
misc_feature      154138..189022
/note="assembly_fragment"
BASE COUNT      55798 a 38604 c 38373 g 54845 t 1402 others
ORIGIN
alignment_scores:
Quality: 43.00 Length: 11
Ratio: 4.778 Gaps: 0
Percent Similarity: 81.818 Percent Identity: 63.636
alignment_block:
US-09-444-281-27 x AC016270 ..
Align seg 1/1 to: AC016270 from: 1 to: 189022
1 HisGluAlaGluProGluAlaGluProIleMet 11
||||| ||||||| |||||||:
43795 CACGAGCCAGACCTGAGCCGAGCCCTGCT 43827
seq_name: gb_in:AMAPID22
seq_documentation_block:
LOCUS      AMAPID22      607 bp      mRNA      INV      03-MAY-1993
DEFINITION A.mellifera Apid22 mRNA.
ACCESSION X72576
VERSION X72576.1 GI:297064
KEYWORDS antibiotic peptide; apid22 gene; Apidaecin; precursor protein.
SOURCE      honeybee.
ORGANISM    Apis mellifera
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Hymenoptera; Apocrita;
Aculeata; Apoidea; Apidae; Apis.
1 (bases 1 to 607)
Casteels-Josson,K.
Direct Submission
Submitted (04-MAR-1993) K. Casteels-Josson, Memorial Sloan
Kettering Cancer Center, 1275 York Avenue, New York 10021, NY, USA
2 (bases 1 to 607)
Casteels-Josson,K., Capaci,T., Casteels,P. and Tempst,P.
Apidaecin multipetide precursor structure: a putative mechanism
for amplification of the insect antibacterial response
EMBO J. 12 (4), 1569-1578 (1993)
932323697
FEATURES
location/Qualifiers
1..607
/oranism="Apis mellifera"
/db_xref="taxon:7460"
1..14
20..76
/gene="Apid22"
/product="secretion signal"
20..454
/gene="Apid22"
/codon_start=1
/product="Apidaecin precursor"
/protein_id="CAA51168.1"
/db_xref="GI:297065"

```


<pre>ELGAVVYVMNMGCGHMADISPIRITCQSDPSMGCISLAVGLDGNOCGFDPINANGAPD</pre>					
<pre>GDCKAFTHHGSSLGIVYKRSQVLEGEAVGIKSFSGTSLSGLDVDGNHTPDLVLGSLAD</pre>					
<pre>TAAIFRARFVNLVSDEIFIDPRADIDEQPCACAGRLVCVDIKICFSYVAAPSVSYSPV</pre>					
<pre>ALDYMDIADTDTRRLRGQVRYRFVLSRGLDDLRRHOSGTWALKQHBRVCDDTYFOLOE</pre>					
<pre>NVKDKRALGVTLVSLGRPPPLRGRAPOGDELPTVPAILNAHQSTORTIEHLFKOQGCG</pre>					
<pre>ODKIOSNNLERYOFCSNISDFEPQALPMLDMGRTALFPALSGOEFHGLEVTYNLP</pre>					
<pre>DPSRRQAOGSDDHENAOVLTVLASLRYSGYRALDSVEKPLCLSNDSASHCELCGNPM</pre>					
<pre>KRGAOVTFLLISTSGITILETELEVKKLLLATISEQLDPSVRAHYTELPUISGSV</pre>					
<pre>ATPOOLFESGEVKGESAMSEKNDVSCSRKYEVVSYNQOSGLNTLGSFLNIMPHEITA</pre>					
<pre>NGKALLYPRVYLEEGOGGKGKIGICSPRNVIQLDVDSDRARRRELGOPEOEPEKV</pre>					
<pre>ESTSMIPPVASEKRNVLTDCQAKCVFESCPLSFDRAAIVLHWGRILMSFTLEE</pre>					
<pre>YMAVTSLEYIVANITYANKSIKNILLRDASTVIPAWVYLDPDAVVYVGVPMMYTILA</pre>					
<pre>VAGLVLYALVLYLLMKCFGRFRNSPSSPTNTHRAHLAVQPSAMEARGCTYGMDS</pre>					
<pre>SRSRPREPCCSTIQ</pre>					
a	258 c	276 g	211 t		
<pre>..</pre>					
<pre>MMYI2380 ..</pre>					
<pre>MMYI2380 from: 1 to: 915</pre>					
<pre>odguAlagluProilemet 11</pre>					
<pre> </pre>					
<pre>GGAGCCGGAGCCGGCTGCTC 378</pre>					
ID73					
ock:	1010 bp	mRNA	INV	01-SEP-1999	
liferA Apid73 mRNA.					
GI:297066					
ic peptide; apid73 gene; Apidaecin; precursor protein.					
lifierA					
a; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;					
a; Neoptera; Endopterygota; Hymenoptera; Apocrita;					
s ; Apoidea; Apidae; Apis.					
s 1 to 1010)					
-Josson,K., Capaci,T., Casteels,P. and Tempst,P.					
n multiplepeptide precursor structure: a putative mechanism					
lication of the insect antibacterial response					
12 (4), 1569-1578 (1993)					
s 1 to 1010)					
-Josson,K.					
ubmission					
ogd (04-MAR-1993) K. Casteels-Josson, Memorial Sloan					
g Cancer Center, 1275 York Avenue, New York 10021, NY, USA					
Location/Qualifiers					
1. 1010					
/organism="Apis mellifera"					
/db_xref="taxon:7450"					
1. 14					
<15. .68					
/gene="Apid73"					
/product="secretion signal"					
<15. .866					
/gene="Apid73"					
/codon_start=1					
/product="Apidaecin precursor"					
/protein_id="CA511169.1"					
/db_xref="GI:4539289"					
/db_xref="SWISS-PROT:Q06602"					
rtranslation="KFALAIIIVTVAHVFGNTLNIDPTRRLRLREARKPEAEFGNNN					
rtvsiIPQPPPHRLREAREPEPNNNRVYIPQPPPHRLRLREXELPAEPNNNR					

VTISPPRPRLRRREAEPEAEPCNNRPVYIPQRPRLRRREAEPEAEPCNNRPVY
ISQRPRLRRREAEPEAEPCNNRPVYIPQRPRLRRREAEPEAEPCNNRPVYIP
QRPRLRRREAEPEAEPCNNRPVYIPQRPRLRRREAEPEAEPCNNRPVYIPQ
RPHPRRI
15. .866
/gene="Apid73"
66. .863
/gene="Apid73"
/product="Apidectin"
misc_signal
866. .870
/note="ATTTA"
misc_signal
902. .906
/note="ATTTA"
polya_signal
906. .911
662. .967
1005. .1010
primer_bind
319 a 279 c 185 g 227 t
BASE COUNT
ORIGIN

alignment_scores:
Quality: 42.00 Length: 8
Ratio: 5.250 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-27 x AMAPID73

Align seg 1/1 to: AMAPID73 from: 1 to: 1010

2 GluA1agluProGluA1agluPro 9
|||||
198 GAGCGTGAACGCGAAGCTGAACCC 221

seq_name: gb_in:DDU66367

seq_documentation_block:
LOCUS DDU66367 1390 bp mRNA INV 28-AUG-1996
DEFINITION Dictyostelium discoideum Sapa (sapa) mRNA, partial cds.
ACCESSION U66367
VERSION U66367.1 GI:1513233
KEYWORDS
SOURCE Dictyostelium discoideum.
ORGANISM Dictyostelium discoideum
REFERENCE 1 (bases 1 to 1390)
AUTHORS Loomis, W.F.
TITLE Direct Submission
JOURNAL Submitted (08-AUG-1996) Dept. of Biology 0322, University of
California, San Diego, 9500 Gilman Drive, La Jolla, CA 92093-032,
USA

FEATURES
source
1. .1390
Location/Qualifiers
/organism="Dictyostelium discoideum"
/strain="AX4"
/db_xref="taxon:44689"
/chromosome="4"
/map="4.053"
/cell_line="AX4"
1. .1372
/gene="sapa"
/note="sapa"
/note="similar to human proactivator polypeptide precursor
sapsin A region, Swiss-Prot Accession Number P07602"
/codon_start=2
/product="Sapa"
/protein_id="AAB06759.1"
/db_xref="GI:1513234"
/translation="LPKEYAODCEYIIINNYGLVRLILNRESPENCIMMELCSKSS
SELIMPINVAVGEVGEICVEYIYFANOTETOTIOLYLDMDCKL
LKSNNWISTCNCNLQIEYEPQIIVAVEGMAPELQCKIKCCSSSSSTNDPFISSSTI

gene
CDS

DEICIFESICAEENLEENKLTLEDIIKVDYDFCKIIPAAVTKDCVAMASNTPIAKM
LEKNSPGVQCKLNCAPPTPTSPSTIKIDVINKKEICTYIGFAEKYGLSALDS
QVYKTFLEONECKRLPGVVDVNCNEVNELEFRAVYQLEKFPPTETICQLFSPTAGSE
SSDASEIKINNNSYIDCOICEVLANVVDKMYTTHFEDKLNKLCSLRSLEPCLN
FVSYGSDSEIQIITVOGESAABACNQIACPTSSSSSYGSASGASGASGASGSGC
SGCYLE"
BASE COUNT 485 a 193 c 225 g 487 t
ORIGIN

alignment_scores:
Quality: 42.00 Length: 8
Ratio: 5.250 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-27 x DDU66367/rev

Align seg 1/1 to reverse of: DDU66367 from: 1 to: 1390

2 GluA1agluProGluA1agluPro 9
|||||
1341 GAGCGAGACCTGAGCGAAGCTT 1318

seq_name: gb_in:DDU72746

seq_documentation_block:
LOCUS DDU72746 1486 bp mRNA INV 19-MAR-1997
DEFINITION Dictyostelium discoideum cysteine proteinase (cprc) mRNA, complete
cds.
ACCESSION U72746
VERSION U72746.1 GI:1644501
KEYWORDS
SOURCE Dictyostelium discoideum.
ORGANISM Dictyostelium discoideum
REFERENCE 1 (bases 1 to 1486)
AUTHORS Ord, T., Adessi, C., Wang, L. and Freeze, H. H.
TITLE The cysteine proteinase gene cprc in Dictyostelium discoideum has a
serine-rich domain that contains GluNAc-1-P
Arch. Biochem. Biophys. 339 (1), 64-72 (1997)
JOURNAL 97223364
MEDLINE 2 (bases 1 to 1486)
REFERENCE Ord, T., Adessi, C., Wang, L. and Freeze, H. H.
AUTHORS Direct Submission
TITLE Submitted (27-SEP-1996) The Burnham Institute, 10901 N. Torrey
Pines Rd. La Jolla, CA 92093, USA
JOURNAL

FEATURES
source
1. .1486
Location/Qualifiers
/organism="Dictyostelium discoideum"
/strain="AX-2"
/db_xref="taxon:44689"
9. .1391
/gene="cprc"
9. .1391
/gene="cprc"
/codon_start=1
/product="cysteine proteinase"
/protein_id="AAC47487.1"
/db_xref="GI:1644502"
/translation="MKVLSALCVLLVAVFAKQDLSEVEYRNAFTNNMIAHORRYSE
EENGKYNIFKAMNDYVNEWNTGSEYVGLVFPADISNEEYRAYLGTTPDASLEMT
ESDKIFPDASAVDRTQGAVPPIKNOGCGGCGFSTTGAEAGVAYLANKKNNVLS
EONLIDCSGYSNGNGCGGLMTLAFEYIINNKGIDTFESSPYTAEDGKKKPNKNA
AOLSSVNVTSGSSPDLAAKVTGQPTVAIDASNOYLYSGIYNRPASSSTYDLDG
VLAVFGTSGSSSGSHGSSQSAAGSSQSAAGSSQSAAGSSQSAAGSSQSAAGSSQ
QSESGSLYSGSYSGSSQSGSGNSGAAVKQGTGASGSGSGSGSGSGSGSSVSGSAG
SASGSASGSSSGSSNSNGVYPTAGDYIVKNSMGTSMGMGYIIMTKGNMNOGCIATM
ASRPTAAVASL"

gene
CDS

BASE COUNT 448 a 314 c 278 g 446 t
ORIGIN

alignment_scores:
Quality: 42.00 Length: 8
Ratio: 5.250 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-27 x DD072746/rev

Align seg 1/1 to reverse of: DD072746 from: 1 to: 1486

2 GIUAAGIUPROGIUAAGIUPRO 9
|||||
1204 GAGCGGAACCTGAGCGGAACCT 1181

THIS PAGE BLANK (USPTO)

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:38:37 : Search time 24.75 Seconds
(without alignments)
10.001 Million cell updates/sec

Title: US-09-444-281-27
Perfect score: 59
Sequence: 1 HEAEEPEEPIW 11

Scoring table:
BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 212252 seqs, 22503292 residues

Total number of hits satisfying chosen parameters: 212252

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
Issued_Patents_AA:*
1: /cgn2_6/prodata/2/1aa/5A_COMB.pep:*
2: /cgn2_6/prodata/2/1aa/5B_COMB.pep:*
3: /cgn2_6/prodata/2/1aa/6A_COMB.pep:*
4: /cgn2_6/prodata/2/1aa/6B_COMB.pep:*
5: /cgn2_6/prodata/2/1aa/PCUTUS_COMB.pep:*
6: /cgn2_6/prodata/2/1aa/backfillseq1.pep:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	39	66.1	486	2 US-08-942-423-3	Sequence 3, Appl1
2	38	64.4	20	1 US-08-208-181A-8	Sequence 8, Appl1
3	38	64.4	20	1 US-08-208-181A-10	Sequence 10, Appl1
4	38	64.4	27	1 US-07-603-782A-1	Sequence 1, Appl1
5	38	64.4	229	2 US-08-394-600B-20	Sequence 20, Appl1
6	38	64.4	229	4 US-08-944-483-30	Sequence 30, Appl1
7	38	64.4	229	5 PCT-US95-02513-20	Sequence 20, Appl1
8	38	64.4	256	2 US-08-230-428B-4	Sequence 4, Appl1
9	37	62.7	226	4 US-08-944-483-41	Sequence 41, Appl1
10	37	62.7	227	4 US-08-944-483-40	Sequence 40, Appl1
11	37	62.7	247	2 US-08-851-974-1	Sequence 1, Appl1
12	37	62.7	247	2 US-08-851-974-4	Sequence 4, Appl1
13	37	62.7	247	2 US-09-213-390-1	Sequence 1, Appl1
14	37	62.7	247	2 US-09-213-390-4	Sequence 4, Appl1
15	37	62.7	248	2 US-08-851-974-3	Sequence 3, Appl1
16	37	62.7	248	2 US-09-213-390-3	Sequence 3, Appl1
17	37	62.7	302	4 US-09-457-046B-18	Sequence 18, Appl1
18	37	62.7	625	3 US-08-996-139-15	Sequence 15, Appl1
19	37	62.7	625	4 US-08-995-659-15	Sequence 15, Appl1
20	37	62.7	625	4 US-09-215-649A-15	Sequence 15, Appl1
21	36	61.0	364	2 US-08-651-940-2	Sequence 2, Appl1
22	36	61.0	364	4 US-09-025-580-37	Sequence 37, Appl1
23	36	61.0	372	4 US-09-286-904-24	Sequence 24, Appl1
24	35	59.3	36	2 US-08-942-423-12	Sequence 12, Appl1
25	35	59.3	433	1 US-08-522-166-7	Sequence 7, Appl1
26	35	59.3	433	1 US-08-488-382A-7	Sequence 7, Appl1
27	35	59.3	433	2 US-08-480-912-7	Sequence 7, Appl1

28	35	59.3	466	2 US-08-836-791-9	Sequence 9, Appl1
29	35	59.3	486	2 US-08-942-423-2	Sequence 2, Appl1
30	35	59.3	2713	5 PCT-US96-01735-1	Sequence 1, Appl1
31	34	57.6	10	1 US-08-468-674B-86	Sequence 86, Appl1
32	34	57.6	10	1 US-08-468-674B-88	Sequence 88, Appl1
33	34	57.6	10	1 US-08-780-571-86	Sequence 86, Appl1
34	34	57.6	10	1 US-08-780-571-88	Sequence 88, Appl1
35	34	57.6	10	4 US-09-012-669F-45	Sequence 45, Appl1
36	34	57.6	13	4 US-08-932-082-6	Sequence 6, Appl1
37	34	57.6	59	1 US-08-306-871-25	Sequence 25, Appl1
38	34	57.6	59	1 US-08-569-959-25	Sequence 25, Appl1
39	34	57.6	65	1 US-08-468-674B-71	Sequence 71, Appl1
40	34	57.6	65	1 US-08-780-571-71	Sequence 71, Appl1
41	34	57.6	88	1 US-08-468-674B-75	Sequence 75, Appl1
42	34	57.6	88	1 US-08-780-571-75	Sequence 75, Appl1
43	34	57.6	326	4 US-09-058-389A-3	Sequence 3, Appl1
44	34	57.6	347	3 US-09-059-369-2	Sequence 2, Appl1
45	34	57.6	456	4 US-09-058-389A-2	Sequence 2, Appl1

ALIGNMENTS

RESULT 1
US-08-942-423-3
; Sequence 3, Application US/08942423
; Patent No. 5891673
GENERAL INFORMATION:
APPLICANT: Hashimoto, Yasuhiro
APPLICANT: Takemoto, Yoshihiro
TITLE OF INVENTION: Lck Binding Protein
NUMBER OF SEQUENCES: 68
CORRESPONDENCE ADDRESS:
ADDRESSEE: Syntex (U.S.A.) Inc.
STREET: 3401 Hillview Ave.
CITY: Palo Alto
STATE: California
COUNTRY: U.S.A.
ZIP: 94303
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/942,423
FILING DATE: 01-Oct-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/362,715
FILING DATE: 23-DEC-1994
ATTORNEY/AGENT INFORMATION:
NAME: Perles, Rohan
REGISTRATION NUMBER: 35,752
REFERENCE/DOCKET NUMBER: 28260
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 852-1698
TELEFAX: (415) 496-3529
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 486 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: linear
MOLECULE TYPE: protein
HYPOTHEICAL: NO
ANTI-SENSE: NO
FRAGMENT TYPE: N-terminal
ORIGINAL SOURCE:
ORGANISM: LCK BINDING PROTEIN
US-08-942-423-3

Query Match 66.1%; Score 39; DB 2; Length 486;
Best Local Similarity 77.8%; Pred. No. 35;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 HEAPEPEEP 9
:|||||:
Db 360 YEAPPEEP 368

RESULT 2

US-08-208-181A-8
; Sequence 8, Application US/08208181A
; Patent No. 5654167
; GENERAL INFORMATION:
; APPLICANT: Gabay, Joelle E.
; APPLICANT: Nathan, Carl F.
; TITLE OF INVENTION: ANTIMICROBIAL PROTEINS, COMPOSITIONS
; TITLE OF INVENTION: CONTAINING SAME AND USES THEREOF
; NUMBER OF SEQUENCES: 23
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Fish & Richardson P.C.
; STREET: 2200 Sand Hill Road, Suite 100
; CITY: Menlo Park
; STATE: CA
; COUNTRY: USA
; ZIP: 94025
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/208,181A
; FILING DATE: 08-MAR-1994
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Bozicevic, Karl
; REGISTRATION NUMBER: 28,807
; REFERENCE/DOCKET NUMBER: 06514/024002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 322-5070
; TELEFAX: (415) 854-0875
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 20 amino acids
; TYPE: amino acid
; STRANDEDNESS: unknown
; TOPOLOGY: unknown
; MOLECULE TYPE: protein
; FRAGMENT TYPE: N-terminal
US-08-208-181A-8

Query Match 64.4%; Score 38; DB 1; Length 20;
Best Local Similarity 54.5%; Pred. No. 1.6;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEPEPIM 11
:|||||:
Db 5 HEAPHSRPM 15

RESULT 3

US-08-208-181A-10
; Sequence 10, Application US/08208181A
; Patent No. 5654167
; GENERAL INFORMATION:
; APPLICANT: Gabay, Joelle E.
; APPLICANT: Nathan, Carl F.
; TITLE OF INVENTION: ANTIMICROBIAL PROTEINS, COMPOSITIONS
; TITLE OF INVENTION: CONTAINING SAME AND USES THEREOF
; NUMBER OF SEQUENCES: 23
; CORRESPONDENCE ADDRESSES:

ADDRESSEE: Fish & Richardson P.C.
STREET: 2200 Sand Hill Road, Suite 100
CITY: Menlo Park
STATE: CA
COUNTRY: USA
ZIP: 94025

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/208,181A
FILING DATE: 08-MAR-1994
CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:
NAME: Bozicevic, Karl
REGISTRATION NUMBER: 28,807
REFERENCE/DOCKET NUMBER: 06514/024002
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 322-5070
TELEFAX: (415) 854-0875
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 20 amino acids
TYPE: amino acid
STRANDEDNESS: unknown
TOPOLOGY: unknown
MOLECULE TYPE: protein
FRAGMENT TYPE: N-terminal
US-08-208-181A-10

Query Match 64.4%; Score 38; DB 1; Length 20;
Best Local Similarity 54.5%; Pred. No. 1.6;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEPEPIM 11
:|||||:
Db 5 HEAPHSRPM 15

US-07-603-782A-1
; Sequence 1, Application US/07603782A
; Patent No. 5200319

GENERAL INFORMATION:
APPLICANT: Arnaout, Amin M.
APPLICANT: McCluskey, Robert T.
APPLICANT: Niles, John L.
TITLE OF INVENTION: Diagnosis of Glomerulonephritis
NUMBER OF SEQUENCES: 1
CORRESPONDENCE ADDRESSES:
ADDRESSEE: Fish & Richardson
STREET: 225 Franklin Street
CITY: Boston
STATE: Massachusetts

COUNTRY: U.S.A.
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb storage
COMPUTER: IBM PS/2 Model 502 or 55SX
OPERATING SYSTEM: IBM P.C. DOS (Version 3.30)
SOFTWARE: WordPerfect (Version 5.0)
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/603,782A
FILING DATE: 19901025
CLASSIFICATION: 436
PRIOR APPLICATION DATA:
PRIOR APPLICATION DATA: including application
PRIOR APPLICATION DATA: described below: 1
APPLICATION NUMBER: 07/428,286
FILING DATE: 27-OCT-1989
ATTORNEY/AGENT INFORMATION:

NAME: Clark, Paul T.
REGISTRATION NUMBER: 30,162
REFERENCE/DOCKET NUMBER: 00786/034002
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 542-5070
TELEFAX: (617) 542-8906
TELEX: 200154
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 27
TYPE: AMINO ACID
TOPOLOGY: linear
US-07-603-782A-1

Query Match 64.4%; Score 38; DB 1; Length 27;
Best Local Similarity 54.5%; Pred. No. 2.3;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPTM 11
|||:|:|
Db 5 HEAPHSRPYM 15

RESULT 5
US-08-394-600B-20
Sequence 20, Application US/08394600B
Patent No. 5843693
GENERAL INFORMATION:
APPLICANT: Halenbeck, Robert F.
APPLICANT: Jewell, David A.
APPLICANT: Kochs, Kirsten E.
APPLICANT: Kriegluer, Michael
APPLICANT: Perez, Carl
TITLE OF INVENTION: Compositions for the inhibition of
TITLE OF INVENTION: Protein Hormone Formation and Uses Thereof
NUMBER OF SEQUENCES: 28
CORRESPONDENCE ADDRESS:
ADDRESSEE: McAndrews, Held & Malloy, Ltd.
STREET: 500 West Madison Street, 34th Floor
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60661
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: EC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/394,600B
FILING DATE: 02/27/95
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Donald J. Pochopien
REGISTRATION NUMBER: 32,167
REFERENCE/DOCKET NUMBER: 820,005/11850US05
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312/707-8889
TELEFAX: 312/707-9155
TELEX:
INFORMATION FOR SEQ ID NO: 20:
SEQUENCE CHARACTERISTICS:
LENGTH: 229 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-394-600B-20

Query Match 64.4%; Score 38; DB 2; Length 229;
Best Local Similarity 54.5%; Pred. No. 23;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPTM 11
|||:|:|
Db 5 HEAPHSRPYM 15

RESULT 6
US-08-944-483-30
Sequence 30, Application US/08944483
Patent No. 6232456
GENERAL INFORMATION:
APPLICANT: COHEN, MAURICE
APPLICANT: COLLITS, TRACEY L.
APPLICANT: FRIEDMAN, PAULA N.
APPLICANT: GRANADOS, EDWARD N.
APPLICANT: KLASS, MICHAEL R.
APPLICANT: RUSSELL, JOHN C.
APPLICANT: STEWART, KENT D.
TITLE OF INVENTION: NOVEL SERINE PROTEASE REAGENTS
TITLE OF INVENTION: OF THE PROSTATE
NUMBER OF SEQUENCES: 76
CORRESPONDENCE ADDRESS:
ADDRESSEE: Abbott Laboratories
STREET: 100 Abbott Park Road
CITY: Abbott Park
STATE: IL
COUNTRY: USA
ZIP: 60064-3500
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM compatible
OPERATING SYSTEM: DOS
SOFTWARE: FASTSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/944,483
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Becker, Cheryl L.
REGISTRATION NUMBER: 35,441
REFERENCE/DOCKET NUMBER: 6183.US.01
TELECOMMUNICATION INFORMATION:
TELEPHONE: 847/935-1729
TELEFAX: 847/938-2623
TELEX:
INFORMATION FOR SEQ ID NO: 30:
SEQUENCE CHARACTERISTICS:
LENGTH: 229 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: NO. 6232456e
US-08-944-483-30

Query Match 64.4%; Score 38; DB 4; Length 229;
Best Local Similarity 54.5%; Pred. No. 23;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPTM 11
|||:|:|
Db 5 HEAPHSRPYM 15

RESULT 7
PCT-US95-02513-20
Sequence 20, Application PC/TUS9502513
GENERAL INFORMATION:

APPLICANT: Halenbeck, Robert F.
APPLICANT: Jewell, David A.
APPLICANT: Koths, Kirston E.
APPLICANT: Kriegluer, Michael
APPLICANT: Perez, Carl
TITLE OF INVENTION: Compositions for the inhibition of
TITLE OF INVENTION: Protein Hormone Formation and Uses thereof
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &
ADDRESSEE: Borun
STREET: 6300 Sears Tower, 233 South Wacker Drive
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60606-6402
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/02513
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Williams Jr., Joseph A.
REGISTRATION NUMBER: 38,659
REFERENCE/DOCKET NUMBER: 27527/32404
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312/474-6900
TELEFAX: 312/474-0448
TELEX: 25-3856
INFORMATION FOR SEQ ID NO: 20:
SEQUENCE CHARACTERISTICS:
LENGTH: 229 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
PCT-US95-02513-20

Query Match 64.4%; Score 38; DB 5; Length 229;
Best Local Similarity 54.5%; Pred. No. 23;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPIM 11
|||:|:|
DB 5 HEAPHSPRYM 15

RESULT 8
US-08-230-428B-4
Sequence 4, Application US/08230428B
Patent No. 5998378
GENERAL INFORMATION:
APPLICANT: Kriegluer, Michael
APPLICANT: Perez, Carl
APPLICANT: Halenbeck, Robert F.
APPLICANT: Jewell, David A.
APPLICANT: Koths, Kirston E.
TITLE OF INVENTION: Compositions For the Inhibition Of TNF
TITLE OF INVENTION: Hormone Formation And Uses Thereof (As Amended)
NUMBER OF SEQUENCES: 14
CORRESPONDENCE ADDRESS:
ADDRESSEE: CHIRON CORPORATION Intellectual Property - R440
STREET: 4560 Horton Street, P.O. Box 8097
CITY: Emeryville
STATE: California
COUNTRY: United States of America
ZIP: 94662-8097
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/230,428B
FILING DATE: 19-APR-1994
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/905,546
FILING DATE: 25-JUN-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/395,253
FILING DATE: 16-AUG-1989
ATTORNEY/AGENT INFORMATION:
NAME: Saveriede, Paul B.
REGISTRATION NUMBER: 36,914
REFERENCE/DOCKET NUMBER: 0820.004
TELECOMMUNICATION INFORMATION:
TELEPHONE: (510) 601-2718
TELEFAX: (510) 655-3542
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 236 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-230-428B-4

Query Match 64.4%; Score 38; DB 2; Length 256;
Best Local Similarity 54.5%; Pred. No. 26;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPIM 11
|||:|:|
DB 32 HEAPHSPRYM 42

RESULT 9
US-08-944-483-41
Sequence 41, Application US/08944483
Patent No. 6232456
GENERAL INFORMATION:
APPLICANT: COHEN, MAURICE
APPLICANT: COLPITTS, TRACEY L.
APPLICANT: FRIEDMAN, PAULA N.
APPLICANT: GRANADOS, EDWARD N.
APPLICANT: KLAAS, MICHAEL R.
APPLICANT: RUSSELL, JOHN C.
APPLICANT: STEWART, KENT D.
APPLICANT: STROUPE, STEVEN D.
TITLE OF INVENTION: NOVEL SERINE PROTEASE REAGENTS
TITLE OF INVENTION: AND METHODS USEFUL FOR DETECTING AND TREATING DISEASES
NUMBER OF SEQUENCES: 76
CORRESPONDENCE ADDRESS:
ADDRESSEE: Abbott Laboratories
STREET: 100 Abbott Park Road
CITY: Abbott Park
STATE: IL
COUNTRY: USA
ZIP: 60064-3500
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/944,483
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER:

FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Becker, Cheryl L.
REGISTRATION NUMBER: 35,441
REFERENCE/DOCKET NUMBER: 6183.US.01
TELECOMMUNICATION INFORMATION:
TELEPHONE: 847/935-1729
TELEFAX: 847/938-2623
TELEX:
INFORMATION FOR SEQ. ID NO: 41:
SEQUENCE CHARACTERISTICS:
LENGTH: 226 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: No. 6232456e
US-08-944-483-41

Query Match 62.7%; Score 37; DB 4; Length 226;
Best Local Similarity 54.5%; Pred. No. 33;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 1 HEAPEAPEPM 11
| | | : | |
DB 5 HEAKPHSRPYM 15

RESULT 10
US-08-944-483-40
Sequence 40, Application US/08944483
Patent No. 6232456
GENERAL INFORMATION:
APPLICANT: COHEN, MAURICE
APPLICANT: COLPITS, TRACEY L.
APPLICANT: FRIEDMAN, PAULA N.
APPLICANT: GRANADOS, EDWARD N.
APPLICANT: KLAS, MICHAEL R.
APPLICANT: RUSSELL, JOHN C.
APPLICANT: STEWART, KENT D.
APPLICANT: STROUPE, STEVEN D.
TITLE OF INVENTION: NOVEL SERINE PROTEASE REAGENTS
TITLE OF INVENTION: AND METHODS USEFUL FOR DETECTING AND TREATING DISEASES
NUMBER OF SEQUENCES: 76
CORRESPONDENCE ADDRESS:
ADDRESSEE: Abbott Laboratories
STREET: 100 Abbott Park Road
CITY: Abbott Park
STATE: IL
COUNTRY: USA
ZIP: 60064-3500
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/944,483
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Becker, Cheryl L.
REGISTRATION NUMBER: 35,441
REFERENCE/DOCKET NUMBER: 6183.US.01
TELECOMMUNICATION INFORMATION:
TELEPHONE: 847/935-1729
TELEFAX: 847/938-2623
TELEX:
INFORMATION FOR SEQ. ID NO: 40:

SEQUENCE CHARACTERISTICS:
LENGTH: 227 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: No. 6232456e
US-08-944-483-40

Query Match 62.7%; Score 37; DB 4; Length 227;
Best Local Similarity 54.5%; Pred. No. 33;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 1 HEAPEAPEPM 11
| | | : | |
DB 5 HEAKPHSRPYM 15

RESULT 11
US-08-851-974-1
Sequence 1, Application US/08851974
Patent No. 5858758
GENERAL INFORMATION:
APPLICANT: Hillman, Jennifer L.
APPLICANT: Corley, Neil C.
APPLICANT: Shah, Puryi
TITLE OF INVENTION: HUMAN SERINE PROTEASE PRECURSOR
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/851,974
FILING DATE: Filed Herewith
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0288 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555
TELEFAX: 415-845-4166
INFORMATION FOR SEQ. ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 247 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: NCBI/NCBI
CLONE: 854243
US-08-851-974-1

Query Match 62.7%; Score 37; DB 2; Length 247;
Best Local Similarity 54.5%; Pred. No. 36;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 1 HEAPEAPEPM 11
| | | : | |
DB 25 HEAKPHSRPYM 35

RESULT 12
US-08-851-974-4
; Sequence 4, Application US/08851974
; Patent No. 5858758
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: HUMAN SERINE PROTEASE PRECURSOR
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FASTSEQ for Windows Version 2.0
; CURRENT APPLICATION DATA:
; FILING DATE: Filed Herewith
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0288 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415-855-0555
; TELEFAX: 415-845-4166
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 247 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: GenBank
; CLONE: 306682
; US-08-851-974-4

Query Match 62.7%; Score 37; DB 2; Length 247;
Best Local Similarity 54.5%; Pred. No. 36;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPRIM 11
|||:|:|
DB 25 HEAKPHSRPYM 35

RESULT 13
US-09-213-390-1
; Sequence 1, Application US/09213390
; Patent No. 5965711
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Corley, Neil C.
; APPLICANT: Shah, Purvi
; TITLE OF INVENTION: HUMAN SERINE PROTEASE PRECURSOR
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA

ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FASTSEQ for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/213,390
; FILING DATE:
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: 08/851,974
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0288 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415-855-0555
; TELEFAX: 415-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 247 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: NCANOT01
; CLONE: 854243
; US-09-213-390-1

Query Match 62.7%; Score 37; DB 2; Length 247;
Best Local Similarity 54.5%; Pred. No. 36;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPRIM 11
|||:|:|
DB 25 HEAKPHSRPYM 35

RESULT 14
US-09-213-390-4
; Sequence 4, Application US/09213390
; Patent No. 5965711
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Corley, Neil C.
; APPLICANT: Shah, Purvi
; TITLE OF INVENTION: HUMAN SERINE PROTEASE PRECURSOR
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FASTSEQ for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/213,390
; FILING DATE:
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: 08/851,974
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0288 US
; TELECOMMUNICATION INFORMATION:

TELEPHONE: 415-855-0555
TELEFAX: 415-845-4166
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 247 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: GenBank
CLONE: 306682
US-09-213-390-4

QY 1 HEAPEAPEPTM 11
Db 25 HEAKPHSRPYM 35
Search completed: January 4, 2002, 08:40:57
Job time: 140 sec

Query Match 62.7%; Score 37; DB 2; Length 247;
Best Local Similarity 54.5%; Pred. No. 36;
Matches 6; Conservat. 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPTM 11
Db 25 HEAKPHSRPYM 35

RESULT 15
US-08-851-974-3
Sequence 3, Application US/08851974
Patent No. 5858758
GENERAL INFORMATION:
APPLICANT: Hillman, Jennifer L.
APPLICANT: Corley, Neil C.
TITLE OF INVENTION: HUMAN SERINE PROTEASE PRECURSOR
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FASTSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/851,974
FILING DATE: Filed Herewith
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0288 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555
TELEFAX: 415-845-4166
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 248 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: GenBank
CLONE: 206690
US-08-851-974-3

Query Match 62.7%; Score 37; DB 2; Length 248;
Best Local Similarity 54.5%; Pred. No. 36;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

THIS PAGE BLANK (USPTO)

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:38:37 ; Search time 53.46 Seconds
(without alignments)
15.241 Million cell updates/sec

Title: US-09-444-281-27
Perfect score: 59
Sequence: 1 HEAEPFAEPIM 11

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 522463 seqs, 74073290 residues
Total number of hits satisfying chosen parameters: 522463

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq_1101:*

1:	/SID58/gcgdata/geneseq/geneseq/AA1980.DAT:*
2:	/SID58/gcgdata/geneseq/geneseq/AA1981.DAT:*
3:	/SID58/gcgdata/geneseq/geneseq/AA1982.DAT:*
4:	/SID58/gcgdata/geneseq/geneseq/AA1983.DAT:*
5:	/SID58/gcgdata/geneseq/geneseq/AA1984.DAT:*
6:	/SID58/gcgdata/geneseq/geneseq/AA1985.DAT:*
7:	/SID58/gcgdata/geneseq/geneseq/AA1986.DAT:*
8:	/SID58/gcgdata/geneseq/geneseq/AA1987.DAT:*
9:	/SID58/gcgdata/geneseq/geneseq/AA1988.DAT:*
10:	/SID58/gcgdata/geneseq/geneseq/AA1989.DAT:*
11:	/SID58/gcgdata/geneseq/geneseq/AA1990.DAT:*
12:	/SID58/gcgdata/geneseq/geneseq/AA1991.DAT:*
13:	/SID58/gcgdata/geneseq/geneseq/AA1992.DAT:*
14:	/SID58/gcgdata/geneseq/geneseq/AA1993.DAT:*
15:	/SID58/gcgdata/geneseq/geneseq/AA1994.DAT:*
16:	/SID58/gcgdata/geneseq/geneseq/AA1995.DAT:*
17:	/SID58/gcgdata/geneseq/geneseq/AA1996.DAT:*
18:	/SID58/gcgdata/geneseq/geneseq/AA1997.DAT:*
19:	/SID58/gcgdata/geneseq/geneseq/AA1998.DAT:*
20:	/SID58/gcgdata/geneseq/geneseq/AA1999.DAT:*
21:	/SID58/gcgdata/geneseq/geneseq/AA2000.DAT:*
22:	/SID58/gcgdata/geneseq/geneseq/AA2001.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and score derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	39	66.1	6797	22	AA831558
2	38	64.4	27	12	AA811996
3	38	64.4	27	14	AA834444
4	38	64.4	237	13	AA820509
5	38	64.4	236	16	AA845403
6	38	64.4	256	16	AA85639
7	37	62.7	149	22	AAU00451
8	37	62.7	173	22	AA872305
9	37	62.7	246	12	AA813253
10	37	62.7	247	19	AA84158
11	37	62.7	261	19	AA854091

12	37	62.7	267	21	AA90291	Human peptidase, H
13	37	62.7	267	22	AA820156	Human protein SECP
14	37	62.7	278	21	AA858142	Lung cancer associ
15	37	62.7	302	22	AAE00235	Protein encoded by
16	37	62.7	353	19	AA863703	Truncated rat RSK3
17	37	62.7	567	20	AA868788	Polyketide fragme
18	37	62.7	568	21	AA85658	Human Acinus S pro
19	37	62.7	571	20	AA841704	Human PRO351 prote
20	37	62.7	571	21	AA844260	Human PRO351 (UNO3
21	37	62.7	571	21	AA824046	Human PRO351 prote
22	37	62.7	583	21	AA85659	Human Acinus S' pr
23	37	62.7	625	19	AA863200	Murine osteoclast
24	37	62.7	625	19	AA869558	Murine NF-KB recep
25	37	62.7	625	19	AA868294	Murine NF-KB recep
26	37	62.7	625	21	AA859509	OBM binding protei
27	37	62.7	625	21	AA833649	A mouse receptor a
28	37	62.7	625	21	AA804427	Murine receptor ac
29	37	62.7	625	22	AAE01994	Murine RANK (recep
30	37	62.7	732	19	AA863715	Rat RSK3 protein.
31	37	62.7	1341	21	AA85657	Human Acinus L pro
32	37	61.0	75	21	AA802199	Human secreted pro
33	36	61.0	108	21	AA833906	Human secreted pro
34	36	61.0	161	21	AA842933	Arabidopsis thalia
35	36	61.0	181	21	AA817997	Arabidopsis thalia
36	36	61.0	181	21	AA847650	Arabidopsis thalia
37	36	61.0	183	21	AA841769	Human OREF ORF1533
38	36	61.0	192	22	AA814154	Human novel protei
39	36	61.0	196	21	AA842932	Arabidopsis thalia
40	36	61.0	213	21	AA843275	Human OREF ORF3039
41	36	61.0	216	21	AA817996	Arabidopsis thalia
42	36	61.0	216	21	AA847649	Arabidopsis thalia
43	36	61.0	252	21	AA815889	Arabidopsis thalia
44	36	61.0	254	21	AA847624	Arabidopsis thalia
45	36	61.0	264	21	AA847623	Arabidopsis thalia

ALIGNMENTS

RESULT 1	
AA831558 standard; Protein: 6797 AA.	
ID	AA831558;
XX	20-APR-2001 (first entry)
XX	Pimaricin biosynthesis associated polyketide synthase polypeptide.
DE	Polyketide synthase; oxidative modification; metabolite; antibiotic;
KW	anticancer; pimaricin.
KW	Streptomyces natalensis.
XX	OS
XX	WO200077222-A1.
PN	21-DEC-2000.
XX	14-JUN-2000; 2000WO-EP06227.
PD	14-JUN-1999; 99EP-0201893.
PR	(STAM) DSM NV.
PA	Martin JF, Aparicio JF, Collina AJ;
XX	WPI; 2001-080693/09.
XX	N-PSDB; AAF24892.
DR	New polynucleotides encoding enzymes involved in the biosynthesis of
XX	pimaricin, useful for modifying the biosynthesis of pimaricin and in
PT	the synthesis of new compounds
PT	
XX	

PS Disclosure; Page 81-101; 116pp; English.
 XX
 CC The present sequence represents a polyketide synthase which is associated
 CC with the biosynthesis of pimarinin. The polyketide synthase polyketide
 CC is useful for the oxidative modification of a methyl group of a suitable
 CC compound, e.g. a bioactive compound including a secondary metabolite,
 CC antibiotics and anticancer agents. Recombinant cells comprising the
 CC gene are useful for the production of pimarinin. The polyketide synthase
 CC polynucleotide may be over expressed in Streptomyces, leading to an
 CC increase in the biosynthesis of pimarinin, as a source of primers for
 CC amplification reaction and as probes.
 XX
 SQ Sequence 6797 AA:
 Query Match 66.1%; Score 39; DB 22; Length 6797;
 Best Local Similarity 77.6%; Pred. No. 1.5e+03;
 Matches 7; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 HEAPEAEP 9
 |||||
 Db 463 heapep 471
 RESULT 2
 AAR1996
 ID AAR1996 standard; peptide; 27 AA.
 XX
 AC AAR1996;
 XX
 DT 26-JUL-1991 (first entry)
 XX
 DE N-terminal of p29 protein.
 XX
 KW Wegener's granulomatosis; monoclonal antibodies; autoantibodies;
 KM glomerulonephritis; serine protease; antigen.
 XX
 OS Homo sapiens.
 XX
 PN W09106572-A.
 XX
 PD 16-MAY-1991.
 XX
 PF 29-OCT-1990; 90MO-US06277.
 XX
 PR 27-OCT-1989; 89US-0428286.
 XX
 PA (GEHO-) GEN HOSPITAL CORP.
 XX
 PI Arnaud M, McCluskey RT, Niles J;
 XX
 DR WPI; 1991-164137/22.
 XX
 PT Purified P29 protein - used to detect auto-antibodies diagnostic
 PT for Wegener's granulomatosis and conditions associated with
 PT glomerulo nephritis
 XX
 PS Claim 1; Page 22; 33pp; English.
 XX
 CC The p29 protein is a 29 kD antigen which was prepd. by affinity
 CC purification from neutrophil acid extract, using 188 monoclonal
 CC antibodies. The purified antigen migrated on SDS-PAGE as three
 CC close bands, with major component at 29 kD under non-reducing
 CC conditions. It reacted with autoantibodies from patients sera
 CC indicating identity between Wegener's granulomatosis autoantigen.
 CC On isofocusing gels it had a pI of 9.2-9.4. It is a novel serine
 CC protease showing homology with leukocyte elastase and cathepsin G.
 CC The protein or monoclonal Abs can be used to test for the presence
 CC of autoantibodies diagnostic for Wegener's granulomatosis. In
 CC combination with myeloperoxidase (and/or Abs against it), p29 Mabs
 CC can also be used to test for autoantibodies associated with pauci
 CC immune necrotizing and/or crescentic glomerulonephritis.
 XX

SQ Sequence 27 AA:
 Query Match 64.4%; Score 38; DB 12; Length 27;
 Best Local Similarity 54.5%; Pred. No. 7.7;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;
 QY 1 HEAPEAEPIM 11
 |||||
 Db 5 heagphsrypym 15
 RESULT 3
 AAR34444
 ID AAR34444 standard; peptide; 27 AA.
 XX
 AC AAR34444;
 XX
 DT 29-JUL-1993 (first entry)
 XX
 DE N-terminus of p29.
 XX
 KW Pauci-immune necrotizing; crescentic glomerulonephritis; neutrophil;
 KM autoantibodies; Wegener's granulomatosis; microscopic polyarteritis
 KW nodosa; primary MCGN.
 XX
 OS Homo sapiens.
 XX
 PN US5200319-A.
 XX
 PD 06-APR-1993.
 XX
 PF 27-OCT-1989; 89US-0428286.
 XX
 PR 27-OCT-1989; 89US-0428286.
 XX
 PR 25-OCT-1990; 90US-0603782.
 XX
 PA (GEHO-) GEN HOSPITAL CORP.
 XX
 PI Arnaud M, McCluskey RT, Niles JL;
 XX
 DR WPI; 1993-133732/16.
 XX
 PT Diagnosis of pauci-immune necrotizing and/or crescentic
 PT glomerulonephritis - using p29 protein isolated from
 PT neutrophil(s), and which binds auto-antibodies present in serum
 PT of patients with Wegener's granulomatosis
 XX
 PS Disclosure; Page 12; 13pp; English.
 XX
 CC The p29 protein can be isolated from neutrophils, has a mol. wt. of
 CC 29 kD (SDS PAGE), binds disopropyl fluorophosphate, has a pI of 9.2-
 CC 9.4, is capable of binding auto antibodies present in the sera of
 CC individuals afflicted with Wegener's granulomatosis and has the N-
 CC terminal sequence shown. The p29 protein can be used for diagnosis
 CC of Wegener's granulomatosis, microscopic polyarteritis nodosa and
 CC primary MCGN.
 XX
 SQ Sequence 27 AA:
 Query Match 64.4%; Score 38; DB 14; Length 27;
 Best Local Similarity 54.5%; Pred. No. 7.7;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;
 QY 1 HEAPEAEPIM 11
 |||||
 Db 5 heagphsrypym 15
 RESULT 4
 AAR20509
 ID AAR20509 standard; protein; 237 AA.

```

XX AC AAR20509:
XX XX
XX DT 06-MAY-1992 (first entry)
XX DE Human proteinase 3.
XX XX
XX KM Serine protease; polymerase chain reaction; PCR; autoantibody.
XX OS
XX FH Homo sapiens.
XX FT Key
XX FT Peptide 1..6
XX FT /label= signal
XX FT /note= "part of signal peptide"
XX FT Peptide 7..8
XX FT /label= propeptide
XX FT Protein 9..237
XX FT /label= proteinase_3
XX PN MO9200378-A.
XX PD 09-JAN-1992.
XX PF 20-JUN-1991; 91MO-1001142.
XX PR 22-JUN-1990; 90DE-4019984.
XX PA (GBFB-) GBF GES BIOTECHN EQ.
XX PI Jenne D, Tschopp J, Ludemann J, Utecht B, Gross LB;
XX DR WPI: 1992-041560/05.
XX DR N-PSDB; AAQ20727.
XX PT DNA sequence coding for proteinase 3 - allows mass prodn. of
XX PT enzyme, with modifications for improved therapy and diagnosis of
XX PT Wegener's granulomatosis
XX PS Disclosure: Fig 3; 21pp; German.
XX CC The CDNA sequence corresponding to the coding region of the
XX CC proteinase 3 gene was obtained by PCR amplification of cDNA from
XX CC cell line U937 (see AAQ22231 and AAQ22232 for PCR primer sequences). An
XX CC amplified product of 784bp was obtained. The product was cut with
XX CC EcoRI and cloned into EcoRI-digested M13mp18/BAP. The complete
XX CC sequence was determined, and the amino acid sequence it encodes was
XX CC deduced from it.
XX SQ Sequence 237 AA:

Query Match 64.4%; Score 38; DB 13; Length 237;
Best Local Similarity 54.5%; Pred. No. 71;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAEPIIM 11
DB 13 headphsrypm 23

```

```

XX OS Homo sapiens.
XX OS MO9400555-A.
XX PN 06-JAN-1994.
XX PD 25-JUN-1993; 93MO-US06120.
XX PF 25-JUN-1992; 92US-0905546.
XX PR 25-JUN-1992; 92US-0905546.
XX PA (CETU ) CETUS ONCOLOGY CORP.
XX PI Halenbeck RF, Jewell DA, Kothe KE, Kriegler M, Perez C;
XX DR WPI: 1994-026195/03.
XX DR N-PSDB; AAQ54498.
XX PT Cpd. which inhibit formation of mature tumour necrosis factor
XX PT from its precursor - identified using TNF convertase, e.g.
XX PT mutein(s), antibodies or peptide phosphonate(s), for preventing
XX PT and treating sepsis, AIDS, auto-immune disease etc.
XX PS Disclosure: Fig 2; 69pp; English.
XX CC PROTNF refers to TNF having a molecular weight of about 26,000,
XX CC which is the pro-hormone form of TNFa. PROTNF is cleaved to a lower
XX CC molecular weight 'mature' form, pref. 17kD, which, in its multimeric
XX CC (usually trimeric) form, is substantially involved in producing life-
XX CC threatening physiological changes associated with sepsis. PROTNF is
XX CC cleaved by convertase. One TNF convertase is serine protease
XX CC proteinase-3, also called PR-3, P-29B or myeloblastin. A suitable
XX CC source of convertase is the HL60 cell line (or extracts, or the
XX CC culture media in which it is grown). The convertase produced by
XX CC HL60 has been sequenced and is identical to the known lymphocyte
XX CC serine protease PR-3 which has other activities unrelated to TNF
XX CC processing.
XX SQ Sequence 256 AA:

Query Match 64.4%; Score 38; DB 15; Length 256;
Best Local Similarity 54.5%; Pred. No. 76;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAEPIIM 11
DB 32 headphsrypm 42

RESULT 6
AAR85639
ID AAR85639 standard; Protein; 256 AA.
XX AAR85639:
XX AC 23-APR-1996 (first entry)
XX DT MY17 preproPR-3.
XX DE MY17 preproPR-3.
XX PR-3; preproPR-3; MY17; human neutrophil protease-3; serine protease;
XX KM tumour necrosis factor alpha; TNFa; TNFa; HL60; MY17; B cell; T cell;
XX KM tumour necrosis factor alpha convertase; cytokine; septic shock;
XX KM rheumatoid arthritis; cachexia; cerebral malaria; graft-host disease;
XX KM ischaemia/reperfusion injury; autoimmune disease; AIDS.
XX OS Homo sapiens.
XX FH Key
XX FH Peptide 1..25
XX FT /note= "leader sequence present only in preproPR-3"
XX FT Peptide 26..27
XX FT /note= "dipeptide present in proPR-3"

```

FT Protein 28..256 /note="mature PR-3"

FT XX MO9524501-A1.

PN XX 14-SEP-1995.

PD XX 02-MAR-1995; 95WO-0502513.

PE XX 28-FEB-1995; 95US-0395456.

PR XX 07-MAR-1994; 94US-0208574.

PR 19-APR-1994; 94US-0230428.

PR 27-FEB-1995; 95US-0394600.

PA (CETUS) CETUS ONCOLOGY CORP.

PI Halenback RF, Jewell DA, Koths KE, Kriegler M, Perez C;

XX N-PSDB; AAT02365.

DR WPI: 1995-328287/42.

XX N-PSDB; AAT02365.

PT Identification of inhibitors of mature TNFalpha prodn. - useful for

PT treatment of septic shock, rheumatoid arthritis, etc..

XX Example 2; Page 82; 96pp; English.

PS This sequence represents the preproPR-3. PR-3 is active recombinant

CC human neutrophil protease-3. PR-3 is a serine protease, and is a tumour

CC necrosis factor alpha (TNFalpha) convertase. The cDNA encoding this

CC sequence was isolated from the HL60 cell clone MY17. The mature PR-3 can

CC be used in the method of the invention for identifying agents that

CC inhibit cleavage of proTNFalpha to mature TNFalpha. In the method,

CC proTNFalpha is incubated with PR-3 (or another TNFalpha convertase), and

CC the cleavage of the proTNFalpha is measured by a colourimetric assay.

CC This is then repeated in the presence of a test compound that is thought

CC to inhibit this process. The results of the two reactions are then

CC compared to determine whether the test compound is an inhibitor. The

CC cleavage inhibitors can be used in the treatment of septic shock,

CC rheumatoid arthritis, cachexia, cerebral malaria, ischaemia/reperfusion

CC injury, graft-host disease, autoimmune diseases, and AIDS. PR-3 can be

CC used to treat unwanted B cell/T cell interaction by contacting it with

CC T cells to cause the release of membrane-bound cytokines.

CC SQ Sequence 256 AA;

QY Query Match 64.4%; Score 38; DB 16; Length 256;

Best Local Similarity 54.5%; Pred. No. 76;

Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

DB 1 HMAPEAPRIM 11

32 heagphsrpyrm 42

RESULT 7

AAU00451

ID AAU00451 standard: Protein; 149 AA.

XX AAU00451;

AC 29-MAY-2001 (first entry)

DT Protein encoded by sugarcane promoter cDNA clones c51, c511 and c512.

XX Sugarcane promoter region; monocotyledonous plant; stem tissue;

KW insecticide; herbicide; disease resistance; improved food content;

KW beta-glucuronidase; GUS; starch biosynthesis; fatty acid biosynthesis;

KW ADP-glucose pyrophosphorylase; sucrose metabolism; clone c51; c511; c512.

OS Saccharum sp.

XX FH- Key Location/Qualifiers

FT Misc-difference 41

FT /note="Encoded by ACG in cDNA clone c51 (AAS01022)"

FT Misc-difference 92

FT /label="Ala, Val

FT /note="Residue 92 is Ala as deduced from cDNA clones

FT c51 and c511 (AAS01022-AAS01023) and Val as

FT deduced from clone c512 (AAS01024)"

FT Misc-difference 103

FT /note="Encoded by TAG in cDNA clone c511 (AAS01023)"

FT Misc-difference 125

FT /label="Val, Ala

FT /note="Residue 125 is Val as deduced from cDNA clone

FT c51 (AAS01022) and Ala as deduced from clones

FT c511 and c512 (AAS01023-AAS01024)"

FT Misc-difference 130

FT /note="Encoded by TCC in cDNA clone c511 (AAS01023)"

FT Misc-difference 135

FT /note="Encoded by GCC in cDNA clone c511 (AAS01023)"

FT WO200118211-A1.

PN 15-MAR-2001.

PD 01-SEP-2000; 2000WO-AU01033.

PE 02-SEP-1999; 99AU-0002625.

PR (UYOU) UNIV QUEENSLAND.

PA Potier B, Birch RG;

PI WPI: 2001-218560/22.

DR N-PSDB; AAS01022, AAS01023, AAS01024.

XX New sugarcane plant promoters for directing expression of heterologous

XX nucleic acids in a constitutive or tissue-specific manner in

XX monocotyledonous plants

XX Claim 40; Fig 15; 107pp; English.

PS The present sequence represents the polypeptide encoded by sugarcane

CC plant promoter cDNA isolated from clones c51, c511 and c512. Clone c51,

CC c511 and c512 promoter cDNAs are 3 of 11 promoter regions of a

CC transcribable DNA sequence isolated from various sugarcane cDNA clones

CC (AAS01021-AAS01031). Also described are 4 promoter regions of specific

CC transcribed DNA sequences (AAS01032-AAS01035). The nucleic acids are

CC useful for producing transgenic plants, having an altered phenotype and

CC for driving expression of a foreign or endogenous DNA sequence, which

CC encode agronomic properties including insecticide, herbicide, disease

CC resistance, stress tolerance and improved food content, or increased

CC yields. The foreign or endogenous DNA sequence may comprise a region

CC transcribed into an antisense RNA or ribozyme that modulates the

CC expression of a corresponding target gene, or it may encode

CC beta-glucuronidase (GUS), luciferase, neomycin phosphotransferase, a

CC product conferring herbicide tolerance, a product affecting starch

CC biosynthesis or modification, ADP-glucose pyrophosphorylase, a product

CC involved in fatty acid biosynthesis, a product conferring insect

CC resistance, a product altering sucrose metabolism or a gene encoding

CC valuable pharmaceuticals, e.g. antibiotics, secondary metabolites or

CC vaccines. The promoters are capable of directing high level expression in

CC many or all cells of a plant, preferentially in stem or meristem tissue

CC or monocotyledonous plants.

CC SQ Sequence 149 AA;

QY Query Match 62.7%; Score 37; DB 22; Length 149;

Best Local Similarity 100.0%; Pred. No. 64;

Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DB 3 AEPEAP 9

69 aepeap 75

RESULT	8	
AA872305		
ID	AA872305	standard; Protein: 173 AA.
XX		
AC	AA872305;	
XX		
DT	16-MAY-2001	(first entry)
XX		
DE	Sunflower glutamic acid rich protein (GRP) amino acid sequence.	
XX		
KW	Defence-related signalling gene; sunflower; neoxanthin cleavage enzyme;	
KM	NES; amino acid permease; AAP; glutamic acid rich protein; GRP;	
KW	pathogen resistance; abscisic acid metabolism.	
XX		
OS	Helianthus annuus.	
XX		
PN	W0200112801-A2.	
XX		
PD	22-FEB-2001.	
XX		
PF	17-AUG-2000; 2000WO-US22961.	
XX		
PR	18-AUG-1999; 99US-0149656.	
PR	23-MAY-2000; 2000US-0206405.	
XX		
PA	(PION-) PIONEER HI-BRED INT INC.	
XX	(CURA-) CURAGEN CORP.	
PI	Bidney DL, Craasta OR, Hu X, Lu G;	
XX		
DR	WPI; 2001-211215/21.	
DR	N-PSDB; AAF77209.	
PT	Novel isolated defence-related signalling gene isolated from sunflower	
PT	encoding neoxanthin cleavage enzyme, amino acid permease or glutamic	
PT	acid-rich protein useful for increasing resistance of plant to a	
XX	pathogen	
PS	Claim 6; Page 107; 135pp; English.	
XX		
CC	This invention relates to defence-related signalling genes isolated from	
CC	the sunflower (<i>Helianthus annuus</i>). The genes encode a neoxanthin cleavage	
CC	enzyme (NCE), an amino acid permease (AAP) and a glutamic acid rich	
CC	protein (GRP). The signalling gene is useful for increasing the	
CC	resistance of a plant to a pathogen such as fungus, virus, bacterium,	
CC	nematode or insect (e.g. European corn borer), preferably	
CC	<i>Scierotinia</i> spp., <i>Phoma</i> spp., or <i>Phomopsis</i> spp., by stably incorporating a	
CC	construct containing the gene into the genome of the plant. The gene is	
CC	useful for regulating gene expression in a plant, in response to a	
CC	stimulus such as infection with a pathogen, damage from a pathogen,	
CC	hydrogen peroxide, jasmonic acid, methyl jasmonate, salicylic acid,	
CC	oxalic acid or expression of a gene encoding oxalic acid oxidase. The	
CC	genes are also useful for stem-preferred regulation of gene expression in	
CC	a plant. The genes are useful in agriculture, particularly in the	
CC	breeding of crop plants with improved agronomic traits, for modifying	
CC	abscisic acid (ABA) metabolism and for modifying amino acid transport and	
CC	content in plants. The present sequence represents the sunflower	
CC	glutamic acid rich protein (GRP) amino acid sequence.	
XX		
Sequence	173 AA;	
XX		

Query March	62.7%	Score 37	DB 22	Length 123
Best Local Similarity	87.5%	Pred. NO	75	
Matches	7	Conservative	0	Mismatches 1
				Indels 0
				Gaps 0
OY	2	EAEPEAP	9	
db	76	epepeap	83	

RESULT	9
AAR13253	
ID	AAR13253 standard; Protein; 246 AA.
XX	
AC	AAR13253;
XX	
DT	11-OCT-1991 (first entry)
XX	
DE	Human Cytotoxic Cell Protease-X.
XX	
RW	hccpx inhibitor; cytotoxic T-lymphocytes; CTL.
XX	
OS	Homo sapiens.
XX	
PN	WO9110685-A.
XX	
PD	25-JUL-1991.
XX	
PF	17-JAN-1991; 91WO-US00340.
XX	
PR	19-JAN-1990; 90US-0467880.
XX	
PA	(SERA-) SERAGEN INC.
XX	
PI	Bleackley RC, Lobe CG, Paetkau VH, James MN, Murphy M;
XX	
DR	WPI; 1991-237989/32.
N-P	N-PsDB; AAO12865, AAO12866.
XX	
PT	DNA vectors, and inhibitors of cytotoxic cell protease - for
PT	treatment of auto-immune diseases e.g. pernicious anaemia,
PT	rheumatoid arthritis, allo-graft rejection etc.
XX	
PS	Claim 8; Fig 9; 62pp: English.
XX	
CC	The hccpx gene was isolated from cytotoxic T-cell lymphocytes using
CC	murine CCP cDNA sequences as probes. The cDNA sequence (i.e. exons
CC	only) was deduced from the genomic sequence. This is the amino acid
CC	sequence deduced from the coding regions. A structural analysis of
CC	the deduced protein was used to design competitive inhibitors of the
CC	protease. See also AAO12862-5 and AAR13254-R13262.
XX	
SO	Sequence 246 AA:
Query Match	62.7%; Score 37; DB 12; Length 246;
Best Local Similarity	54.5%; Pred.No.1.le+02;
Matches 6; conservative	2; Mismatches 3; Indels 0; Gaps 0;
OY	1 HEAEPPEAEPIIM 11 : : 25 heakphsrpyim 35

RESULT	10
AAW84158	
ID	AAW84158 standard; Protein: 247 AA.
XX	
XX	
AC	AAW84158;
XX	
DT	25-FEB-1999 (first entry)
XX	
DE	A human serine protease precursor (HSP) protein.
XX	
KW	Human serine protease precursor; HSP; Incyte clone 854243;
KW	immunological disorder; cancer; antagonist; allergy; asthma;
KW	Crohn's disease; multiple sclerosis; rheumatoid arthritis; glioma;
KW	lymphoma; myeloma.
XX	
OS	Homo sapiens.
XX	
FH	key
FT	Modified-site 71 Location/Qualifiers

FT Modified-site /note= "potential N-linked glycosylation site"
 FT 104 /note= "potential N-linked glycosylation site"
 FT Disulfide-bond 49 /note= "potential disulphide bridging site"
 FT Disulfide-bond 65 /note= "potential disulphide bridging site"
 FT Disulfide-bond 142 /note= "potential disulphide bridging site"
 FT Disulfide-bond 173 /note= "potential disulphide bridging site"
 FT Disulfide-bond 188 /note= "potential disulphide bridging site"
 FT Disulfide-bond 209 /note= "potential disulphide bridging site"
 FT Disulfide-bond 230 /note= "potential disulphide bridging site"
 FT Disulfide-bond 230 /note= "potential disulphide bridging site"
 FT Region 37-48 /note= "potential substrate binding region"

PN W09850424-A2.

PD 12-NOV-1998.

PF 06-MAY-1998; 98MO-US09096.

PR 07-MAY-1997; 97US-0851974.

PA (INCY-) INCYTE PHARM INC.

PI Corley NC, Hillman JL, Shah P;

DR WPI; 1999-034707/03.

DR N-PSDB; AAV82707.

XX New human serine protease precursor and related nucleic acid.
 PT vectors - for treatment, prevention and diagnosis of immunological
 PT disorders and cancer

PS Claim 1: Fig 1A-C; 56pp; English.

XX The present sequence represents a human serine protease precursor (HSP)
 CC protein. Nucleic acids encoding HSP were first identified in incyte
 CC clone 854243 from the ganglioma tissue cDNA library. The protein
 CC has homology with the rat natural killer cell protease-1 precursor
 CC (RNPR-1) and a human serine protease from cytotoxic T lymphocytes (SECT).
 CC As increased levels of the protein are associated with immunological
 CC disorders and cancers, antagonists are used to treat or prevent
 CC conditions such as allergy, asthma, Crohn's disease, multiple sclerosis,
 CC rheumatoid arthritis, glioma, lymphoma, myeloma etc. The HSP nucleic
 CC acid sequence and its fragments are used as antisense/ribozyme
 CC therapeutics, for detecting and quantifying gene expression (as probes
 CC and primers in standard hybridisation and amplification assays), and for
 CC isolating related sequences and for chromosome mapping.

XX Sequence 247 AA;

Query Match 62.7%; Score 37; DB 20; Length 247;
 Best Local Similarity 54.5%; Pred. No. 1.1e+02;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 1 HEAEPEAPIM 11
 DB 25 heakphsrypm 35

RESULT 11
 AAM54091
 ID AAM54091 standard; Protein; 261 AA.
 AC AAM54091;
 XX

DT 28-SEP-1998 (first entry)
 XX Homo sapiens BE2 sequence.
 DE BARD1; ring protein; BRCA1; breast cancer; risk; diagnosis.
 KW Homo sapiens.
 XX

FT Key Location/Qualifiers

FT Misc-difference 104 /label= Ala, Ser, Pro, Thr

FT Misc-difference 114 /label= Gly

FT Misc-difference 218 /label= Ala

PN W09812327-A2.

PD 26-MAR-1998.

PF 19-SEP-1997; 97WO-US16842.

PR 04-APR-1997; 97US-0042985.

PR 20-SEP-1996; 96US-0025296.

PR 03-APR-1997; 97US-0042611.

PA (TEXA) UNIV TEXAS SYSTEM.

PI Baer R, Bowcock AM;

DR WPI; 1998-230317/20.

DR N-PSDB; AAV24126.

XX DNA sequence encoding BARD1, B123, BE2, BE14, BE31 or BE445 - which
 PT as breast cancer antigen, BRCA1, binding proteins are useful to
 PT identify patient having or at risk of developing cancer

XX Disclosure: Page 272-273; 348pp; English.

XX The sequence is that of a protein which can be used in the
 CC preparation of the recombinant breast cancer antigen, BRCA1, binding
 CC proteins BARD1, B123, BE2, BE14, BE31 or BE445, or a composition for the
 CC detection of a BARD1, B123, BE2, BE14, BE31 or BE445 nucleic acid
 CC sequence, specifically a wild type BARD1 composition for the detection
 CC or purification of BRCA1, useful to identify a patient having, or at
 CC risk of developing cancer. BARD1 can be used in the preparation of an
 CC anti-BARD1 antibody, and in the detection and purification of a BRCA1
 CC protein. BARD1, B123, BE2, BE14, BE31 or BE445 can be used in the
 CC identification of a binding protein agonist or antagonist that alters
 CC the binding of BARD1, B123, BE2, BE14, BE31 or BE445 to BRCA1 or the
 CC biological activity of the BRCA1-BARD1, B123, BE2, BE14, BE31 or BE445
 CC complex. The antibodies can be used to detect BARD1, B123, BE2, BE14,
 CC BE31 or BE445, a specific anti-BARD1 antibody can be used to identify
 CC a patient having or at risk of developing cancer.

XX Sequence 261 AA;

Query Match 62.7%; Score 37; DB 19; Length 261;
 Best Local Similarity 60.0%; Pred. No. 1.1e+02;
 Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

OY 2 EAPEAPIM 11
 DB 219 eaqpeapal 228

RESULT 12
 AAY90291
 ID AAY90291 standard; Protein; 267 AA.
 AC AAY90291;
 XX

DT		24-OCr-2000	(first entry)
XX			
DE		Human peptidase, HPEP-8 protein sequence.	
KW		Human; peptidase; cell proliferative disorder; arteriosclerosis;	
KW		psoriasis; myelofibrosis; cancer; autoimmune disorder; Crohn's disease;	
KW		Inflammatory disorder; AIDS; anaemia; allergy; asthma; arteriosclerosis;	
KW		Grave's disease; multiple sclerosis; scleroderma; infection; diabetes;	
KW		metabolic disorder; Addison's disease; cystic fibrosis; diagnosis;	
KW		glycogen storage disease; obesity; therapy; HPEP-8.	
XX			
OS		Homo sapiens.	
PN		WO200042201-A2.	
PD			
XX		20-JUL-2000.	
PF			
XX		11-JAN-2000; 2000WO-US00641.	
PR		11-JAN-1999; 99US-0172247.	
PR		03-MAY-1999; 99US-0132253.	
PR		27-MAY-1999; 99US-0136653.	
XX			
PA		(INCY-) INCYTE PHARM INC.	
PI			
PI		Bandman O, Hillman JL, Tang YT, Azimzai Y, Baughn MR, Lal P;	
PI		Yue H, Lu DAM;	
DR		WP1: 2000-482832/42.	
DR		N-PSDB: AAA37664.	
PT			
PT		An isolated polypeptide for diagnosis, prevention and treatment of	
PT		cell proliferative, autoimmune/inflammatory and metabolic disorders	
PT		comprises a sequence encoding a human peptidase -	
XX			
PS		Claim 2; Page 99; 131pp; English.	
XX			
CC		This sequence represents a human peptidase, designated HPEP-8. The	
CC		invention relates to 18 human peptidases designated HPEP-1 to HPEP-18,	
CC		respectively. The peptidases can be used for treating a disease or	
CC		condition associated with decreased expression or over expression of	
CC		functional human peptidases. The diseases that can be diagnosed,	
CC		prevented and treated include cell proliferative disorders (such as	
CC		arteriosclerosis, psoriasis, myelofibrosis, and cancers),	
CC		autoimmune/inflammatory disorders (such as AIDS, anaemia, allergies,	
CC		Crohn's disease, asthma, arteriosclerosis, Grave's disease, multiple	
CC		sclerosis, and scleroderma), infections, and metabolic disorders (such as	
CC		Addison's disease, diabetes, cystic fibrosis, glycogen storage diseases	
CC		and obesity).	
XX			
SQ		Sequence 267 AA:	
OY		2 EAEFEAEP 9	
		I I I I I I I I	
Db		247 epepeaeap 254	
Matches	7;	Conservative 0;	Mismatches 1;
Indels			Gaps 0;
Score		Best Local Similarity 62.7%;	Score 37;
Pred. No.		1.2e+02;	Length 267;
Result		AAB20156	ID AAB20156 standard; Protein; 267 AA.
AC		AAB20156;	
DT		30-Apr-2001	(first entry)
XX			
XX		Human protein SECP2.	
XX			
XX		SECP2; secreted protein; human; diagnosis; therapy.	

XX	Homo sapiens.
OS	
XX	WO200105971-A2.
PN	
PD	25-JAN-2001.
XX	
PE	20-JUL-2000; 2000WO-US19890.
XX	
PR	20-JUL-1999; 99US-0144722.
PR	29-NOV-1999; 99US-0167785.
PR	19-JUL-2000; 2000US-0619252.
XX	
PA	(CURA-) CURAGEN CORP.
XX	
PI	Shimkets RA, Fernandes E;
XX	
DR	WPI; 2001-091973/10.
DR	N-PSDB; AAF30189.
XX	
PT	New polypeptide designated SECP, its encoding nucleic acid and its immunospecific antibody, useful for diagnosing, preventing and treating SECP-associated disorders such as cancer .
PT	
XX	
PS	Claim 1; Fig 2; 124pp; English.
CC	
CC	The present sequence is that of novel human protein SECP2,
CC	which is predicted to localise in the macrobody (peroxisome),
CC	and which does not appear to include a signal peptide. The
CC	protein shows homology to human PRO31 protein, and to a region
CC	of human prostatic precursor. The invention provides 9 novel
CC	SECP proteins (see AAB20155-63), nucleic acids encoding them
CC	(see AAF30188-96), antibodies, mutants or fragments. These can
CC	be used to detect, treat or prevent an SECP-associated disorder,
CC	to screen for predisposition to such a disorder, and to identify
CC	agents that modulate the expression or activity of SECP.
XX	
SQ	Sequence 267 AA:
Query Match 62.7%; Score 37; DB 22; Length 267;	
Best Local Similarity 87.5%; Pred.No.1.2e+02;	
Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;	
OY 2 FAEPFAP 9	
	I I I I I I I
Db 247 epeapeap 254	
RESULT 14	
AAB58142	
ID	AAB58142 standard; Protein; 278 AA.
XX	
AC	AAB58142;
XX	
DT	14-MAR-2001 (first entry)
XX	
DE	Lung cancer associated polypeptide sequence SEQ ID 480.
XX	
KM	Human; lung cancer associated protein; neuroprotective; cytostatic;
KM	cardioactive; immunomodulatory; muscular active; vulnerary;
KM	gastrointestinal; nephrotoxic; antineoplastic; gynecological;
KM	antibacterial; diagnosis; neural disorder; immune disorder; reproductive;
KM	proliferative disorder; wound healing; infectious disease.
XX	
OS	Homo sapiens.
XX	
PN	WO200055180-AZ.
XX	
PD	21-SEP-2000.
XX	
PF	08-MAR-2000; 2000WO-US05918.
XX	

```

BR 12-MAR-1999; 99US-0124270.
XX
XX (HOMA-) HUMAN GENOME SCI INC.
PA (ROSE/) ROSEN C A.
XX
XX Ruben SM;
PI WPI: 2000-587514/55.
DR N-PSDB; AAF18018.
XX
XX Lung cancer associated gene sequences, referred to as lung cancer
PT antigens, useful for treatment, prevention, and diagnosis of disorders
PT such as lung cancer -
XX
XX Claim 11; Page 961-962; 1425pp; English.
XX
XX Polynucleotide sequences AAF17982 - AAF18424 encode human lung cancer
CC associated proteins represented in AAB58106 - AAB58548. Lung cancer
CC associated proteins and polynucleotide sequences, their agonists, and
CC antagonists may have neuroprotective; cytostatic; cardioactive;
CC immunomodulatory; muscular active general; vulnery; gastrointestinal
CC general; nephrotoxic; antineoplastic; gynecological; or antibacterial
CC activity. The invention also includes antibodies specific for the
CC protein or polynucleotide sequences. The lung cancer associated
CC polynucleotide sequences may be used for detection of lung cancer,
CC chromosome identification, as chromosome markers, and for numerous other
CC diagnostic or research purposes. The proteins may be used to treat
CC disorders such as neural, immune, muscular, reproductive,
CC gastrointestinal, pulmonary, cardiovascular, renal, and proliferative
CC disorders. The proteins may also be used in the treatment of wounds and
CC infectious diseases. Polynucleotide sequences AAF18425 - AAF18433 and
CC peptide AAB58549 are used in the course of the invention for the
CC identification and characterization of the polynucleotide and protein
CC sequences.
XX
XX Sequence 278 AA:
SQ

```

```

Query Match 62.7%; Score 37; DB 21; Length 278;
Best Local Similarity 54.5%; Pred. No. 1.2e+02;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

```

```

OY 1 HEAPEAEPIIM 11
   |||:|:|
DB 57 heakhsrpyim 67

```

```

RESULT 15
AAE00235
ID AAE00235 standard; Protein; 302 AA.
XX
XX AAE00235;
AC
XX 13-JUN-2001 (first entry)
DT
XX Protein encoded by Probe 9.
DE
XX
XX Transacylase; taxol; paclitaxel biosynthesis; taxoid; probe;
KW transgenic organism; TAX9.
XX
XX Taxus cuspidata.
OS
XX
XX Key Location/Qualifiers
FH Misc-difference 119
FT /note= "Encoded by TCG"
FT Misc-difference 164
FT /note= "Encoded by ATG"
FT Misc-difference 187
FT /note= "Encoded by TGC"
XX
XX WO200123586-A2.
PN
PD -05-APR-2001.

```

```

XX
XX 29-SEP-2000; 2000WO-US27006.
PE
XX
XX 30-SEP-1999; 99US-0411145.
PR 07-DEC-1999; 99US-0457046.
XX
XX (UNIW ) UNIV WASHINGTON STATE RES FOUND.
XX
XX Croteau RB, Walker KD, Schoendorf A, Wildung MR;
PI WPI: 2001-245004/25.
DR N-PSDB; AAD03341.
XX
XX New transacylase enzymes, useful for the high yield production of
PT Taxol(TM), related taxoids and useful intermediates in the in the
PT paclitaxel biosynthetic pathway -
XX
XX Claim 1; Page 108-109; 162pp; English.
XX
XX The present sequence is the amino acid sequence of probe 9.
CC Probe 9 is derived from AT-FOR2 and AT-REV1 primers and is
CC used for screening taxus cuspidata TAX9 full length cDNA clone. The
CC probes are useful for the identification of (nucleic acid sequences
CC encoding) transacylases. The probes isolated from the Taxus genus
CC are useful for the synthetic production of Taxol(TM) and related taxoids,
CC as well as intermediates in the paclitaxel biosynthetic pathway. They
CC can also be used for the creation of transgenic organisms that either
CC produce the transacylases for subsequent in vitro use, or produce the
CC transacylases in vivo. The (nucleic acids encoding) transacylases are
CC also useful for creating specific binding agents that recognise the
CC corresponding transacylases. Binding agents include (fragments of)
CC antibodies or any other agent capable of specifically binding to the
CC groups on the proteins.
XX
XX Sequence 302 AA:
SQ

```

```

Query Match 62.7%; Score 37; DB 22; Length 302;
Best Local Similarity 70.0%; Pred. No. 1.3e+02;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

```

```

OY 2 EAPEAEPIIM 11
   ||:|:|
DB 110 eakpslepiim 119

```

```

Search completed: January 4, 2002, 08:40:26
Job time: 109 sec

```


LOCUS A2518927 390 bp DNA GSS 16-OCT-2000
 DEFINITION RPCI-11-67B18. TV RPCI-11 Homo sapiens genomic clone RPCI-11-67B18.
 ACCESSION A2518927
 VERSION A2518927.1 GI:10829921
 KEYWORDS GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 390)
 Zhao, S., Adams, M.D., Nieman, W., Malek, J., de Jong, P. and Venter, J.C.
 TITLE BAC end sequences of library RPCI-11
 JOURNAL Unpublished (1997)
 COMMENT Other GSSs: RPCI11-67B18.TJ
 Contact: Shaying Zhao
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: szhao@tigr.org
 Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong (pieterdejong.med.bufileo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.bufileo.edu/ordering) or from Research Genet cs (info@resgen.com). BAC end search page: http://www.tigr.org/cdb/humgen/bac_end_search.html. This BAC end was generated during the Rad process and may have higher chance of clone tracking errors.
 Seg primer: T7
 Class: BAC ends.
 FEATURES
 source Location/Qualifiers
 1..390
 /organism="Homo sapiens"
 /db_xref="GDB:7525385"
 /db_xref="taxon:9606"
 /clone="RPCI-11-67B18"
 /clone_lib="RPCI-11"
 /sex="Male"
 /cell_type="Lymphocytes"
 /note="Vector: pBACE3.6; Site_1: EcoRI; Site_2: EcoRI; RPCI11 Human Male BAC Library"
 BASE COUNT 59 a 159 c 84 g 87 t 1 others
 ORIGIN
 alignment_scores:
 Quality: 69.00 Length: 12
 Ratio: 6.273 Gaps: 0
 Percent Similarity: 91.667 Percent Identity: 83.333
 alignment_block:
 US-09-444-281-35 x A2518927/rev ..
 Align seg 1/1 to reverse of: A2518927 from: 1 to: 390
 2 LeuLysLysTrpProTrpTrpProTPaGAGLys 13
 |||||:|||| |||||:|||||||:|||||||:|||||||
 189 TTACAAATAATCCCTGCTGGCCCTGAGAGAGAG 154
 seq_name: gb_est1:BE237369
 seq_documentation_block:
 LOCUS BE237369 415 bp mRNA EST 25-APR-2001
 DEFINITION 146629 MARC 4BOV Bos taurus cDNA 5', mRNA sequence.
 ACCESSION BE237369
 VERSION BE237369.1 GI:9022087
 KEYWORDS EST.
 SOURCE cow.
 ORGANISM Bos taurus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE
 AUTHORS
 TITLE
 JOURNAL
 MEDLINE
 COMMENT
 Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae; Bovidae; Bovinae; Bos.
 1 (bases 1 to 415)
 Smith, T.P.L., Grose, W.M., Freking, B.A., Roberts, A.J., Stone, R.T., Casas, E., Wray, J.E., White, J., Cho, J., Fahrenkrug, S.C., Bennett, G.L., Heaton, M.P., Laegreid, W.W., Rohrer, G.A., Chitko-McKown, C.G., Perlea, G., Holt, L., Karameycheva, S., Liang, F., Quackenbush, J. and Keeler, J.W.
 Sequence evaluation of four pooled-tissue normalized bovine cDNA libraries and construction of a gene index for cattle genome Res. 11 (4), 626-630 (2001)
 Contact: Smith TPL
 USDA, ARS, US Meat Animal Research Center
 PO Box 166, Clay Center, NE 68933-0166, USA
 Tel: 402 762 4366
 Fax: 402 762 4390
 Email: smith@email.marc.usda.gov
 Single pass sequencing. Bases called and alt-trimmed with phred v0.980904.e. Vector identified by cross-match with the -minscore 18 and -mismatch 12 options.
 PCR Primers
 FORWARD: AGGAACACGCTATGACCAT
 BACKWARD: GTTTCCTCAGTACAGACG
 Plate: 47 row: J column: 10
 Seq primer: ATTATGGTGACATATAC.
 FEATURES
 source Location/Qualifiers
 1..415
 /organism="Bos taurus"
 /db_xref="taxon:9913"
 /clone_lib="MARC 4BOV"
 /tissue_type="pooled"
 /lab_host="DH10B"
 /note="Vector: pCMV SPORT6; Site_1: XbaI; Site_2: XhoI; Library made from PMV SV40 tissue from day 20 and day 40 embryos."
 BASE COUNT 106 a 97 c 120 g 91 t 1 others
 ORIGIN
 alignment_scores:
 Quality: 68.00 Length: 8
 Ratio: 8.500 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000
 alignment_block:
 US-09-444-281-35 x BE237369 ..
 Align seg 1/1 to: BE237369 from: 1 to: 415
 4 LysTrpProTrpTrpProTrpPArg 11
 |||||:|||||:|||||:|||||:|||||:|||||
 280 AATGCGCATGCTGGCCCTTGGCGC 303
 seq_name: gb_est1:A0089922
 seq_documentation_block:
 LOCUS A0089922 446 bp mRNA EST 19-APR-2000
 DEFINITION A0089922 Hordeum vulgare subsp. vulgare upper three leaves at heading stage Hordeum vulgare subsp. vulgare upper three leaves at haruna_lib1_121, mRNA sequence.
 ACCESSION A0089922
 VERSION A0089922.1 GI:7613350
 KEYWORDS EST.
 SOURCE Hordeum vulgare subsp. vulgare.
 ORGANISM Hordeum vulgare subsp. vulgare.
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Poideae; Triticeae; Hordeum.
 1 (bases 1 to 446)
 Sato, K., Takahashi, H. and Takeda, K.
 Hordeum vulgare subsp. vulgare cDNA clone
 Unpublished (2000)
 JOURNAL

COMMENT

Contact: Kazuhiro Sato
Research Institute for Bioresources
Okayama University, Barley Germplasm Center
Chuo 2-20-1, Kurashiki, Okayama 710-0046, Japan
Email: kazuhiro@rib.okayama-u.ac.jp,
URL: http://www.rib.okayama-u.ac.jp/barley/
Location/Qualifiers

FEATURES

source

1. 446
/organism="Hordeum vulgare subsp. vulgare"
/cultivar="Haruna Nijo"
/db_xref="taxon:112509"
/clone_lib="haruna_lib1_121"
/clone_lib="Hordeum vulgare subsp. vulgare Upper three
leaves at heading stage"
/tissue_type="Upper three leaves at heading stage"
89 a 130 c 149 g 76 t 2 others

BASE COUNT
ORIGIN

alignment_scores:

Quality: 68.00 Length: 8
Ratio: 8.500 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-35 x AU089922/rev ..

Align seg 1/1 to reverse of: AU089922 from: 1 to: 446

5 TrpProtTrpProtTrpParGarg 12
|||||
188 TGGCGCTGGCTGGCGCGCGCA 165

seq_name: gb_est1:AU198144

seq_documentation_block:

LOCUS AU198144 446 bp mRNA EST 12-JUL-2001
DEFINITION AU198144 Rice green shoot Oryza sativa cDNA clone S15951, mRNA
sequence.
ACCESSION AU198144
VERSION AU198144.1 GI:14714211
KEYWORDS EST.
SOURCE Oryza sativa.
ORGANISM Oryza sativa.
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Ethnarioidae; Oryzaceae; Oryza.
1 (bases 1 to 446)
Sasaki, T. and Yamamoto, K.
Rice cDNA from green shoot (2001)
Unpublished (2001)
Contact: Takuji Sasaki
National Institute of Agrobiological Resources
Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki
305-8602, Japan
Tel: 81-298-38-7441
Fax: 81-298-38-7468
Email: tsasaki@abr.affrc.go.jp, URL: http://rgp.dna.affrc.go.jp/
PROJECT = 'RGP'.
S15951_97A.

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

Location/Qualifiers
1. 446
/organism="Oryza sativa"
/strain="Nipponbare"
/db_xref="taxon:4530"
/clone="S15951"
/clone_lib="Rice green shoot"
/note="Green shoot (8 days old)"
PROJECT = 'RGP'.
S15951_97A.

BASE COUNT

ORIGIN

82 a 153 c 161 g 50 t

alignment_scores:

Quality: 68.00 Length: 11
Ratio: 6.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:

US-09-444-281-35 x AU198144/rev ..

Align seg 1/1 to reverse of: AU198144 from: 1 to: 446

3 LysLysTrpProtTrpProtTrpParGargLys 13
:::|||||
351 CGCGCGCTGGCTGGCGCGCGCGCGG 319

seq_name: gb_est1:AU198162

seq_documentation_block:

LOCUS AU198162 448 bp mRNA EST 12-JUL-2001
DEFINITION AU198162 Rice green shoot Oryza sativa cDNA clone S16019, mRNA
sequence.
ACCESSION AU198162
VERSION AU198162.1 GI:14714231
KEYWORDS EST.
SOURCE Oryza sativa.
ORGANISM Oryza sativa.
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Ethnarioidae; Oryzaceae; Oryza.
1 (bases 1 to 448)
Sasaki, T. and Yamamoto, K.
Rice cDNA from green shoot (2001)
Unpublished (2001)
Contact: Takuji Sasaki
National Institute of Agrobiological Resources
Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki
305-8602, Japan
Tel: 81-298-38-7441
Fax: 81-298-38-7468
Email: tsasaki@abr.affrc.go.jp, URL: http://rgp.dna.affrc.go.jp/
PROJECT = 'RGP'.
S16019_97A.

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

FEATURES

source

1. 448
/organism="Oryza sativa"
/strain="Nipponbare"
/db_xref="taxon:4530"
/clone="S16019"
/clone_lib="Rice green shoot"
/note="Green shoot (8 days old)"
BASE COUNT 85 a 146 c 160 g 57 t
ORIGIN

alignment_scores:

Quality: 68.00 Length: 11
Ratio: 6.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:

US-09-444-281-35 x AU198162/rev ..

Align seg 1/1 to reverse of: AU198162 from: 1 to: 448

3 LysLysTrpProtTrpProtTrpParGargLys 13
:::|||||
314 CGCGCGCTGGCTGGCGCGCGCGCGG 282

seq_name: gb_est1:AU089934

seq_documentation_block:

LOCUS AU089934 475 bp mRNA EST 19-APR-2000
DEFINITION AU089934 Hordeum vulgare subsp. vulgare upper three leaves at
heading stage Hordeum vulgare subsp. vulgare cDNA clone
haruna_lib1_134, mRNA sequence.

```

ACCESSION      AUT089934
VERSION        AUT089934.1
KEYWORDS       EST
SOURCE         Hordeum vulgare subsp. vulgare.
ORGANISM       Hordeum vulgare subsp. vulgare
REFERENCE      Sato, K., Takahashi, H. and Takeda, K.
AUTHORS        Unpublished (2000)
TITLE          Hordeum vulgare subsp. vulgare cDNA clone
JOURNAL        Unpublished (2000)
COMMENT        Contact: Kazuhiko Sato
                Research Institute for Bioreources
                Okayama University, Barley Germplasm Center
                Chuo 2-20-1, Kurashiki, Okayama 710-0046, Japan
                Email: kzasato@rib.okayama-u.ac.jp,
                URL:http://www.rib.okayama-u.ac.jp/barley/.
FEATURES       location/Qualifiers
               1..475
               /organism="Hordeum vulgare subsp. vulgare"
               /cultivar="Hartuna Nijo"
               /db_xref="taxon:112509"
               /clone="hartuna libl 134"
               /clone_lib="Hordeum vulgare subsp. vulgare Upper three
               leaves at heading stage"
               /tissue-type="Upper three leaves at heading stage"
BASE COUNT     100 a 137 c 155 g 83 t
ORIGIN
alignment_scores:
    Quality:   68.00           Length:   8
    Ratio:     8.500           Gaps:     0
Percent Similarity: 100.000    Percent Identity: 100.000
alignment_block:
US-09-444-281-35 x AUT089934/rev ..
Align seg 1/1 to reverse of: AUT089934 from: 1 to: 475
5 TrpproTPTPrpDTPArGArg 12
|||||
194 TGCCCGTGGTGGCCGfGGCGGCCGA 171
seq_name: gb_est1:AUT082117
seq_documentation_block:
LOCUS          AUT082117             578 bp            mRNA            EST            04-FEB-2000
DEFINITION    AUT082117 Rice panicle at ripening stage Oryza sativa cDNA clone
EIL6111, mRNA sequence.
VERSION       AUT082117
KEYWORDS      AUT082117.1 GI:6727452
SOURCE        EST.
ORGANISM      Oryza sativa.
              Oryza sativa
              Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
              Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
              Ehrhartoideae; Oryzaceae; Oryza.
              1 (bases 1 to 578)
              Sasaki,T. and Yamamoto,K.
              Rice cDNA from panicle at ripening stage (2000)
              Unpublished (2000)
              Contact: Takuji Sasaki
              National Institute of Agrobiological Resources
              Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki
              305-8602, Japan
              Tel: 81-298-38-7441
              Fax: 81-298-38-7468
              Email: tsasaki@abr.affrc.go.jp, URL:http://rgp.dna.affrc.go.jp/
              PROJECT = 'RGP'.
FEATURES       Location/Qualifiers
               1..578

```

```

/organism="Oryza sativa"
/strain="Nipponbare"
/db_xref="taxon:4530"
/clone="E11611"
/clone_lib="Rice panicle at ripening stage"
/dev_stage="ripening stage"
/note="Organ: panicle; Rice cDNA from panicle at ripening stage"

```

BASE COUNT 129 a 180 c 179 g 90 t

ORIGIN

alignment_scores:

Quality: 68.00 Length: 11

Ratio: 6.800 Gaps: 0

Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:

US-09-444-281-35 x AU082117/rev ..

Align seg 1/1 to reverse of: AU082117 from: 1 to: 578

3 LysLysTrpProTTrpTTrpArgLys 13
 ::::::::::::::::::::

348 CGCCGCTGCCTGCTGCTGCCTGCAGCGGCGG 316

seq_name: gb_est1:AU198258

seq_documentation_block:

LOCUS AU198258 352 bp mRNA EST 12-Jul-2001

DEFINITION AU198258 Rice green shoot Oryza sativa cDNA clone S16389, mRNA

sequence.

ACCESSION AU198258

VERSION AU198258.1 GI:14711435

KEYWORDS EST.

SOURCE

ORGANISM

Oryza sativa.

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Ehrhartoideae; Oryzaceae; Oryza.

1 (bases 1 to 352)

Sasaki,T. and Yamamoto,K.
 Rice cDNA from green shoot (2001)
 Unpublished (2001)
 Contact: Takuji Sasaki
 National Institute of Agrobiological Resources
 Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki 305-8602, Japan
 Tel: 81-298-38-7441
 Fax: 81-298-38-7468
 Email: tsasakileabr.affrc.go.jp, URL:http://rtp.dna.affrc.go.jp/PROJECT/'RGP'.

S16389_96Z.

FEATURES

Source

1..352 Location/Qualifiers

/organism="Oryza sativa"
 /strain="Nipponbare"
 /db_xref="taxon:4530"
 /clone="S16389"
 /clone_lib="Rice green shoot"
 /note="Green shoot (8 days old)"

BASE COUNT 95 a 95 c 104 g 51 t 7 others

ORIGIN

alignment_scores:

Quality: 66.00 Length: 10

Ratio: 7.333 Gaps: 0

Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:

US-09-444-281-35 x AU198258/rev ..

KEYWORDS
SOURCE house mouse.
ORGANISM Mus musculus
REFERENCE
AUTHORS 1 (bases 1 to 600)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duvall,B., Hamil,C., Islam,H., Longacre,S., Mahmood,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Stokes,R., Tinney,A., von Niederhausen,A. and Wright,D., Weiss,R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
JOURNAL
COMMENT Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert length: 10000 Std Error: 0.00
Plate: 0118 row: 0 column: 21
Seq primer: CACACAGGAACACGCTATGACC
Class: plasmid ends
High quality sequence stop: 600.
Location/Qualifiers
1. 600
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUCG1M0118021"
/clone_lib="Mouse 10kb plasmid UUCG1M library"
/sex="Male"
/lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: pMD42ny; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD42 (g11473211419b/AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 173 a 108 c 181 g 138 t
ORIGIN

alignment_scores:
Quality: 60.00 Length: 9
Ratio: 7.333 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:
US-09-444-281-35 x A2368740/rev ..
Align seg 1/1 to reverse of: A2368740 from: 1 to: 600

2 LysLysLysTrpProTrpTrpProTrp 10
|||||:|||||:|||||:|||||:|||||
384 TTACGGAGGTGGCCATGCTGCTGG 358

seq_name: gb_est2:BE977776

seq documentation block:
LOCUS BE977776 255 bp mRNA EST 04-OCT-2000
DEFINITION bs68h10.y1 Drosophila melanogaster adult testis library Drosophila
melanogaster cDNA clone bs68h10 5', mRNA sequence.
ACCESSION BE977776
VERSION BE977776.1 GI:10608588
KEYWORDS EST.
SOURCE fruit fly.
ORGANISM Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 255)
REFERENCE Andrews, J., Bouffard, G. and Oliver, B.
AUTHORS Drosophila melanogaster testis expressed sequence tags
JOURNAL Unpublished (1999)
COMMENT Contact: Brian Oliver
Laboratory of Cellular and Developmental Biology
NIDDK, National Institutes of Health
6 Center Drive MSC 2715, Bldg 6, Rm BI-13, Bethesda, MD 20892 USA
Fax: (301) 496 5239
Email: oliver@helix.nih.gov
http://www.nidk.nih.gov/intram/people/boliver.htm
Tissue isolation and library construction performed at the National
Institute of Diabetes and Digestive and Kidney Diseases, NIH (see
http://www.nidk.nih.gov/intram/people/boliver.htm). DNA sequencing
and analyses performed by National Institutes of Health Intramural
Sequencing Center (NISC; see http://www.nisc.nih.gov).
Plate: 68 row: h column: 10
Seq primer: M13Rp1 reverse primer (ABI).
Location/Qualifiers
1. 255
/organism="Drosophila melanogaster"
/strain="y1 w[67c1]/Y"
/db_xref="taxon:7227"
/clone="bs68h10"
/clone_lib="Drosophila melanogaster adult testis library"
/sex="male"
/dev_stage="1-5 day adult"
/lab_host="SOLR (Stratagene)"
/note="Organ: testis; Vector: pBluescript SK (Stratagene);
Site_1: Bcor I; Site_2: Xho I; Testes dissected from 1-5
day adult y1 w[67c1]/Y males raised at 25°C. RNA
isolated using Trizol (Life Technologies) and a single
round of Poly(A)+ selection using Oligotex (Qiagen). cDNA
library constructed using Stratagene ZAP-cDNA synthesis
kit. Oligo dT-primed, size fractionated -1-6 kb, and
directionally cloned at EcoRI and XhoI in Uni-ZAP XR.
Following a single round of amplification pBluescript SK
phagemids were mass excised. A distribution channel for
clones is being sought, but not currently available.
Requests for clones cannot be honored."

BASE COUNT 47 a 60 c 101 g 47 t
ORIGIN

alignment_scores:
Quality: 65.00 Length: 15
Ratio: 5.909 Gaps: 1
Percent Similarity: 73.333 Percent Identity: 60.000

alignment_block:
US-09-444-281-35 x BE977776 ..
Align seg 1/1 to: BE977776 from: 1 to: 255

3 LysLysTrpPro.....TrpTrpProTrpArgArgLys 13
|||||:|||||:|||||:|||||:|||||
185 AAGAGCTGCCCCGTGAGGAAAAGTGTGCTGCTTGAGAGAAAGAA 229

seq_name: gb_est1:AV641634

```

seq_documentation_block: 371 bp, mRNA, EST, 15-DEC-2000
LOCUS AV641634 Chlamydomonas reinhardtii 5% CO2 Chlamydomonas reinhardtii
DEFINITION AV641634 Chlamydomonas reinhardtii 5% CO2 Chlamydomonas reinhardtii
ACCESSION AV641634
VERSION AV641634.1 GI:10784962
KEYWORDS
SOURCE EST.
ORGANISM Chlamydomonas reinhardtii.
Chlamydomonas reinhardtii Chlorophyta: Chlorophyceae; Volvocales;
Eukaryota; Viridiplantae; Chlamydomonadales.
REFERENCE 1 (bases 1 to 371)
AUTHORS Asanizu,E., Miura,K., Kuchio,K., Inoue,Y., Fukuzawa,H., Ohyama,K.,
Nakamura,Y. and Tabata,S.
TITLE Generation of expressed sequence tags from low-CO2 and high-CO2
adapted cells of Chlamydomonas reinhardtii
JOURNAL DNA Res. 7 (5), 305-307 (2000)
MEDLINE 20539644
COMMENT Contact: Erika Asanizu
The First Laboratory for Plant Gene Research
Kazusa DNA Research Institute
Yana 1532-3, Kisarazu, Chiba 292-0812, Japan
Email: asanizu@kazusa.or.jp, URL: http://www.kazusa.or.jp/en/plant/.
Location/Qualifiers
1. 371
/organism="Chlamydomonas reinhardtii"
/strain="C9"
/db_xref="taxon:3055"
/clone="HCL037h03_r"
/clone_lib="Chlamydomonas reinhardtii 5% CO2"
/note="Vector: pBluescriptII SK-; Site: 1: EcoRI; Site 2:
XhoI: The cDNA library was constructed from cells cultured
in a medium with bubbling air containing 5% carbon
dioxide"
BASE COUNT 73 a 106 c 141 g 51 t
ORIGIN

alignment_scores:
Quality: 65.00 Length: 10
Ratio: 7.222 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:
US-09-444-281-35 x AV641634 ...
Align seg 1/1 to: AV641634 from: 1 to: 371
3 LysLysTrpProTrpTrpProTrpArgArg 12
+++++|||||
241 GAACGGTGGCGGTGGTGGCGGCGG 270

seq_name: gb_est1:BE129188

seq_documentation_block: 386 bp, mRNA, EST, 21-JUN-2000
LOCUS BE129188
DEFINITION 894021E12.Y1 C. reinhardtii CC-1690, normalized, Lambda Zap II
ACCESSION BE129188
VERSION BE129188.1 GI:8576551
KEYWORDS
SOURCE EST.
ORGANISM Chlamydomonas reinhardtii.
Chlamydomonas reinhardtii Chlorophyta: Chlorophyceae; Volvocales;
Eukaryota; Viridiplantae; Chlamydomonadales.
REFERENCE 1 (bases 1 to 386)
AUTHORS Grossman,A., Davies,J., Federlel,N., Harris,E., Lefebvre,P.,
McDermott,J.P., Silflow,C., Stern,D. and Surzycki,R.
TITLE Analyses of the Chlamydomonas reinhardtii Genome: A Model,
Unicellular System for Analyzing Gene Function and Regulation in
Vascular Plants; project phase 2
JOURNAL Unpublished (2000)
COMMENT Contact: Elizabeth H. Harris

```

```

DCMB Box 91000
Duke University
Durham, NC 27708-1000, USA
Tel: 919 613 8164
Fax: 919 613 8177
Email: chlamydeduke.edu.
Location/Qualifiers
1. 386
/organism="Chlamydomonas reinhardtii"
/strain="CC-1690 wild type ml+ 21gr"
/db_xref="taxon:3055"
/clone_lib="C. reinhardtii CC-1690, normalized, Lambda Zap
II"
/note="Vector: pBluescript II SK-; Site 1: EcoRI; Site 2:
XhoI: This library, constructed by John Davies and Jeffrey
McDermott, combines cDNAs from CC-1690 cells grown to
mid-log phase in TAP (acetate-containing) medium in the
light, TAP medium in the dark, HS (minimal) medium in
ambient levels of CO2 and HS medium bubbled with 5% CO2.
polyA mRNA was purified from each sample, pooled and cDNA
synthesized. The cDNA was directionally cloned into lambda
Zap II (Stratagene) in the EcoRI (5') and XhoI (3') sites.
pBluescript II SK- plasmids were excised from the lambda
Zap clones by superinfection with Exsist (Stratagene)
phage. The library was normalized using method 4 described
in Bonaldo et al (1996) Genome Research 6: 791-806."
BASE COUNT 79 a 109 c 142 g 56 t
ORIGIN

alignment_scores:
Quality: 65.00 Length: 10
Ratio: 7.222 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:
US-09-444-281-35 x BE129188 ..
Align seg 1/1 to: BE129188 from: 1 to: 386
3 LysLysTrpProTrpTrpProTrpArgArg 12
+++++|||||
246 GAACGGTGGCGGTGGTGGCGGCGG 275

```

THIS PAGE BLANK (USPTO)

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:47:23 : Search time 50.17 seconds
(without alignments)
37,902 Million cell updates/sec

Title: US-09-444-281-35
Perfect score: 91
Sequence: 1 ILKKPMWPMWRK 13

Scoring table: BLOSUM62
Gapop 10.0, Gapext 0.5

Searched: 473505 seqs, 146272329 residues
Total number of hits satisfying chosen parameters: 473505

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: SPTEMBL_17:*
2: sp_archaea:*
3: sp_bacteria:*
4: sp_fungi:*
5: sp_human:*
6: sp_invertebrate:*
7: sp_mammal:*
8: sp_mhc:*
9: sp_organelle:*
10: sp_plant:*
11: sp_rodent:*
12: sp_virus:*
13: sp_vertebrate:*
14: sp_unclassified:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	57	62.6	723	12	09DUC4
2	54	59.3	1173	12	0990M4
3	54	59.3	1173	12	0990M3
4	54	59.3	1173	12	0990M2
5	54	59.3	1173	12	0990M1
6	54	59.3	1383	12	084712
7	52	57.1	746	12	094H31
8	52	57.1	1018	12	094H33
9	51	56.0	399	4	094H31
10	50	54.9	327	10	094H33
11	49	53.8	148	5	026590
12	49	53.8	298	1	094H33
13	49	53.8	467	5	019573
14	49	53.8	528	5	026589
15	49	53.8	528	5	094H33
16	49	53.8	725	12	0990M4
17	49	53.8	802	5	094H33
18	49	53.8	1245	3	094H33
19	49	53.8	1940	5	002456

20	48	52.7	49	12	09D80	09d80 tt virus. o
21	48	52.7	111	5	018753	018753 caenorhabd
22	48	52.7	428	11	09JMG0	09JMG0 mus musculu
23	48	52.7	431	11	099ML4	099ML4 mus musculu
24	48	52.7	748	12	09D781	09d81 tt virus. o
25	47.5	52.2	114	2	09X8C2	09x8c2 streptomyce
26	47	51.6	141	11	09CZAI	09czai mus musculu
27	47	51.6	165	10	09SNN3	09snn3 oryza sativ
28	47	51.6	504	2	P96143	P96143 thermoactin
29	46.5	51.1	352	2	P73417	P73417 synochocyst
30	46.5	51.1	620	12	091H07	091H07 avian infec
31	46.5	51.1	621	12	066196	066196 avian infec
32	46.5	51.1	621	12	091H14	091H14 avian infec
33	46.5	51.1	621	12	091H13	091H13 avian infec
34	46.5	51.1	621	12	091H11	091H11 avian infec
35	46.5	51.1	621	12	091H10	091H10 avian infec
36	46.5	51.1	621	12	091H15	091H15 avian infec
37	46.5	51.1	621	12	091H12	091H12 avian infec
38	46.5	51.1	621	12	091H09	091H09 avian infec
39	46.5	51.1	621	12	091H08	091H08 avian infec
40	46.5	51.1	621	12	0991L8	0991L8 avian infec
41	46.5	51.1	621	12	0991L7	0991L7 avian infec
42	46.5	51.1	621	12	0991L6	0991L6 avian infec
43	46.5	51.1	625	12	09QCP6	09qcp6 avian infec
44	46.5	51.1	630	12	066197	066197 avian infec
45	46.5	51.1	630	12	098WQ0	098wq0 avian infec

ALIGNMENTS

RESULT 1
ID 09DUC4 PRELIMINARY; PRT; 723 AA.
AC 09DUC4;
DT 01-MAR-2001 (TEMBLrel. 16, Created)
DT 01-MAR-2001 (TEMBLrel. 16, Last sequence update)
DT 01-JUN-2001 (TEMBLrel. 17, Last annotation update)
DE ORF1.
OS TP VIRUS.
OC Viruses; ssDNA viruses; unclassified ssDNA viruses.
OX NCBI_TaxID=68887;
RN [1]
RC SEQUENCE FROM N.A.
RA Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=MF-TTV9;
RX PubMed-11080484;
RA Okamoto H., Nishizawa T., Tawara A., Peng Y., Takahashi M.,
RA Kishimoto J., Tanaka T., Miyakawa Y., Mayumi M.;
RT "Species-specific TP viruses in humans and nonhuman primates and their
RL phylogenetic relatedness.";
RL Virology 277:368-378(2000).
DR EMBL: AB041959; BAB19313.1;
DR InterPro: IPR001563; Serine_carpept.
DR PROSITE: PS00131; CARBOXYPEPT_SER_SER; UNKNOWN_1.
SO SEQUENCE 723 AA; 85393 MW; 232D003098766344 CRC64;

Query Match 62.68; Score 57; DB 12; Length 723;
Best Local Similarity 100.0%; Pred. No. 4.1;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 6 PWMPPRR 12
| | | | |
Db 2 PWMPPRR 8

RESULT 2
0990M4

ID 0990M4 PRELIMINARY; PRT; 1173 AA.
AC 0990M4;
DT 01-JUN-2001 (TREMBlrel. 17, Created)
DE 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DE SPIKE GLYCOPROTEIN.
GN S.
OS Human coronavirus (strain 229E).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=11137;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=229E;
RA Bonavia A., Holmes K.V.;
RT "Viral and cellular changes in a human cell line persistently infected with human coronavirus HCoV-229E."
RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF344186; AAK32188.1;
SQ SEQUENCE 1173 AA; 128669 MW; ABC6E0A75EBDBA4 CRC64;

Query Match 59.3%; Score 54; DB 12; Length 1173;
Best Local Similarity 85.7%; Pred. No. 16;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 4 KWPMPW 10
Db 1113 KWPMPW 1119

RESULT 3
0990M3 PRELIMINARY; PRT; 1173 AA.
AC 0990M3;
DT 01-JUN-2001 (TREMBlrel. 17, Created)
DE 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DE SPIKE GLYCOPROTEIN.
GN S.
OS Human coronavirus (strain 229E).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=11137;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=229E;
RA Bonavia A., Holmes K.V.;
RT "Viral and cellular changes in a human cell line persistently infected with human coronavirus HCoV-229E."
RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF344187; AAK32189.1;
SQ SEQUENCE 1173 AA; 128683 MW; 9E236816082A81A CRC64;

Query Match 59.3%; Score 54; DB 12; Length 1173;
Best Local Similarity 85.7%; Pred. No. 16;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 4 KWPMPW 10
Db 1113 KWPMPW 1119

RESULT 4
0990M2 PRELIMINARY; PRT; 1173 AA.
AC 0990M2;
DT 01-JUN-2001 (TREMBlrel. 17, Created)
DE 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DE SPIKE GLYCOPROTEIN.
GN S.

OS Human coronavirus (strain 229E).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=11137;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=229E;
RA Bonavia A., Holmes K.V.;
RT "Viral and cellular changes in a human cell line persistently infected with human coronavirus HCoV-229E."
RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF344188; AAK32190.1;
SQ SEQUENCE 1173 AA; 128653 MW; 8B658FCBBD1842DA CRC64;

Query Match 59.3%; Score 54; DB 12; Length 1173;
Best Local Similarity 85.7%; Pred. No. 16;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 4 KWPMPW 10
Db 1113 KWPMPW 1119

RESULT 5
0990M1 PRELIMINARY; PRT; 1173 AA.
AC 0990M1;
DT 01-JUN-2001 (TREMBlrel. 17, Created)
DE 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DE SPIKE GLYCOPROTEIN.
GN S.
OS Human coronavirus (strain 229E).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=11137;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=229E;
RA Bonavia A., Holmes K.V.;
RT "Viral and cellular changes in a human cell line persistently infected with human coronavirus HCoV-229E."
RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF344189; AAK32191.1;
SQ SEQUENCE 1173 AA; 128760 MW; B73A165A6270152A CRC64;

Query Match 59.3%; Score 54; DB 12; Length 1173;
Best Local Similarity 85.7%; Pred. No. 16;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 4 KWPMPW 10
Db 1113 KWPMPW 1119

RESULT 6
084712 PRELIMINARY; PRT; 1383 AA.
AC 084712;
DT 01-NOV-1996 (TREMBlrel. 01, Created)
DE 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
DE SPIKE PROTEIN.
OS Porcine epidemic diarrhea virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=28295;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BRI/87;
MEDLINE=94231173; PubMed=8176382;

RA Duarte M., Laude H.;
RT "Sequence of the spike protein of the porcine epidemic diarrhoea virus."
RL J. Gen. Virol. 75:1195-1200(1994).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-BRI/87;
RX MEDLINE=93389433; PubMed=8397280;
RA Bridgen A., Duarte M., Tobler K., Laude H., Ackermann M.;
RT "Sequence determination of the nucleocapsid protein gene of the porcine epidemic diarrhoea virus confirms that this virus is a coronavirus related to human coronavirus 229E and porcine transmissible gastroenteritis virus."
RL J. Gen. Virol. 74:1795-1804(1993).
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN-BRI/87;
RX MEDLINE=94120721; PubMed=8291230;
RA Duarte M., Tobler K., Bridgen A., Rasschaert D., Ackermann M., Laude H.;
RT "Sequence analysis of the porcine epidemic diarrhoea virus genome between the nucleocapsid and spike protein genes reveals a polymorphic ORF."
RL Virology 198:466-476(1994).
DR EMBL: Z25483; CAA80971.1; -;
DR InterPro: IPR002551; Corona_S1.
DR pfam: PF01600; Corona_S1; 1.
DR pfam: PF01601; Corona_S2; 1.
RT CONFLICT 422 422 Y -> N (IN REF. 1).
SQ SEQUENCE 1383 AA; 151404 MW; 741C84D5DD3BCAD CRC64;

Query Match 59.3%; Score 54; DB 12; Length 1383;
Best Local Similarity 85.7%; Pred. No. 18;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 KMPMPW 10
Db 1322 KMPMPW 1328

RESULT 7
O9JH31 PRELIMINARY; PRT; 746 AA.
AC O9JH31;
DT 01-OCT-2000 (TRENBLREL. 15, Created)
DT 01-OCT-2000 (TRENBLREL. 15, Last sequence update)
DE ORF1.
OS TT virus.
OC Viruses; ssDNA viruses; unclassified ssDNA viruses.
OX NCBI_TaxID=68887;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-TJN02;
RA Okamoto H.;
RL Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-TJN02;
RA Ukiha M., Okamoto H., Nishizawa T., Tawara A., Takahashi M., Iizuka H., Miyakawa Y., Mayumi M.;
RT "The entire nucleotide sequences of two distinct TT virus (TTV) isolates (TJN01 and TJN02) remotely related to the original TTV isolates."
RL Arch. Virol. 0:0-0(2000).
DR EMBL: AB028669; BAA94878.1; -;
SQ SEQUENCE 746 AA; 88561 MW; E0B22953AE76AE3E CRC64;

Query Match 57.1%; Score 52; DB 12; Length 746;
Best Local Similarity 66.7%; Pred. No. 19;

Matches 6; Conservative 1; Mismatches 2; Indels 0; Gaps 0;
Qy 5 WPMWPRK 13
Db 3 WPMWPRR 11

RESULT 8
O9HKX3 PRELIMINARY; PRT; 1018 AA.
AC O9HKX3;
DT 01-MAR-2001 (TRENBLREL. 16, Created)
DT 01-MAR-2001 (TRENBLREL. 16, Last sequence update)
DT 01-JUN-2001 (TRENBLREL. 17, Last annotation update)
DE CONSERVED HYPOTHETICAL MEMBRANE PROTEIN.
GN TA0470.
OS Thermoplasma acidophilum.
OC Archaea; Euryarchaeota; Thermoplasmatales; Thermoplasmataceae;
OX Thermoplasma.
RN NCBI_TaxID=2303;
RP SEQUENCE FROM N.A.
RC STRAIN-DSM 1728;
RX MEDLINE=20479972; PubMed=11029001;
RA Ruepp A., Graml W., Santos-Martinez M.-L., Koretke K.K., Volker C., Mewes H.-W., Fishman D., Stocker S., Lupas A.N., Baumister W.;
RT "The genome sequence of the thermophilic scavenger Thermoplasma acidophilum."
RL Nature 407:508-513(2000).
DR EMBL: AL445064; CAC11612.1; -;
DR InterPro: IPR000731; HMGCR_Patched_5TM.
DR PROSITE: PS50156; SSD; 1.
KW Complete proteome.
SQ SEQUENCE 1018 AA; 112323 MW; 83BE84D3C74B853 CRC64;

Query Match 57.1%; Score 52; DB 1; Length 1018;
Best Local Similarity 66.7%; Pred. No. 25;
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TLKMPWP 9
Db 1004 LMKKMPWP 1012

RESULT 9
O9Y4N1 PRELIMINARY; PRT; 299 AA.
AC O9Y4N1;
DT 01-NOV-1999 (TRENBLREL. 12, Created)
DT 01-MAY-2000 (TRENBLREL. 13, Last sequence update)
DT 01-MAY-2000 (TRENBLREL. 13, Last annotation update)
DE HYPOTHETICAL 34.0 KDA PROTEIN (FRAGMENT).
GN DKFZP434C192.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=TESTIS;
RA Ansoorge W., Winkner U., Mewes H.W., Gassenhuber J., Wiemann S.;
RL Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL: AL096753; CAB46428.2; -;
KW Hypothetical protein.
FT NON_TER 1
SQ SEQUENCE 299 AA; 34032 MW; 6B8DB6DE6A88239A CRC64;

Query Match 56.0%; Score 51; DB 4; Length 299;
Best Local Similarity 85.7%; Pred. No. 11;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 6 PWMWR 12
 ||||| 1
 Db 37 PWMWR 43

RESULT 10
 ID 09AUN3 PRELIMINARY; PRT: 327 AA.
 AC 09AUN3:
 DT 01-JUN-2001 (TREMBLrel. 17, Created)
 DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
 DE 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
 DE HYPOTHEITICAL PROTEIN.
 OS Oryza sativa (Rice).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 OC Eriactoidae; Oryzae; Oryza.
 OX NCBI_TaxID=4530;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Spiegel L.A., King L., Kirchoff K.A., de la Bastide M., Preston R.R.,
 RA Nascimiento L.U., Vil M.D., Baker J.P., Miller B., Cunnius D.M.,
 RA Kuit K.H., Rodriguez S., Santos L., Zutavern T., Ballja V.S.,
 RA Shah R.S., Bahret A., Bai H.P., O'Shaughnessy A., Dedhia N.N.,
 RA McCombie W.R.;
 RT "Genomic Sequence For Oryza sativa, Nipponbare Strain, Chromosome X,
 RT Clone OSUBA0058B19, Complete Sequence."
 RL Submitted (MAR-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AC083945; AAK13143.1;
 SQ SEQUENCE 327 AA; 36672 MW; 5CCA9080664BD0CA CRC64;

Query Match 54.9%; Score 50; DB 10; Length 327;
 Best Local Similarity 100.0%; Pred. No. 17;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 7 WWPWR 12
 ||||| 1
 Db 119 WWPWR 124

RESULT 11
 ID 026590 PRELIMINARY; PRT: 148 AA.
 AC 026590:
 DT 01-NOV-1996 (TREMBLrel. 01, Created)
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
 DT 01-OCT-2000 (TREMBLrel. 15, Last annotation update)
 DE MYOSIN HEAVY CHAIN (FRAGMENT).
 OS Schistosoma mansoni (Blood fluke).
 OC Eukaryota; Metazoa; Platyhelminthes; Rhabditophora; Neodermata;
 OC Trematoda; Digenea; Strigeidida; Schistosomatidae; Schistosomatidae;
 OC Schistosoma
 OX NCBI_TaxID=6183;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Schmitz J., Symmons P., Dargatz H., Kunz W.;
 RL Submitted (FEB-1992) to the EMBL/GenBank/DBJ databases.
 DR EMBL; M81338; AAA29906.1;
 KW Myosin.
 FT NON_TER
 SQ SEQUENCE 148 AA; 17923 MW; C7EDA5A0BBE14DDA CRC64;

Query Match 53.8%; Score 49; DB 5; Length 148;
 Best Local Similarity 62.5%; Pred. No. 11;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
 QY 1 LKKRPMW 8
 :||| 1
 Db 30 VLKRPWM 37

RESULT 12
 ID 09Y806 PRELIMINARY; PRT: 298 AA.
 AC 09Y806:
 DT 01-NOV-1999 (TREMBLrel. 12, Created)
 DT 01-NOV-1999 (TREMBLrel. 12, Last sequence update)
 DE 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
 DE HYPOTHEITICAL 33.7 KDA PROTEIN APE2577.
 GN APE2577.
 OS Aeropyrum pernix.
 OC Archaea; Crenarchaeota; Desulfurococcales; Desulfurococaceae;
 OC Aeropyrum.
 OX NCBI_TaxID=56636;
 RN [1]
 RP SEQUENCE FROM N.A.

RC STRAIN=K1
 RX MEDLINE=99310339; PubMed=10382966;
 RA Kawarabayashi Y., Hino Y., Horikawa H., Yamazaki S., Halkawa Y.,
 RA Jin-no K., Takahashi M., Sekine M., Baba S.-I., Aikai A., Kosugi H.,
 RA Hosoyama A., Fukui S., Nagai Y., Nishijima K., Nakazawa H.,
 RA Takamiya M., Masuda S., Funahashi T., Tanaka T., Kudoh Y.,
 RA Yamazaki J., Kushiida N., Oguchi A., Aoki K.-I., Kubota K.,
 RA Nakamura Y., Nomura N., Sako Y., Kikuchi H.;
 RT "Complete genome sequence of an aerobic hyper-thermophilic
 RT crenarchaeon, Aeropyrum pernix K1."
 RL DNA Res. 6:83-101(1999).
 DR EMBL; AP000064; BAA81594.1;
 DR InterPro: IPR002787; DUF85.
 DR Pfam: PF01932; DUF85; 1.
 KW Hypothetical protein: Complete proteome.
 SQ SEQUENCE 298 AA; 33666 MW; FCB9C6EC93FE231 CRC64;

Query Match 53.8%; Score 49; DB 1; Length 298;
 Best Local Similarity 60.0%; Pred. No. 21;
 Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
 QY 2 LKKRPMW 11
 :||| 1
 Db 102 LKRPWMWR 111

RESULT 13
 ID 019573 PRELIMINARY; PRT: 467 AA.
 AC 019573:
 DT 01-NOV-1996 (TREMBLrel. 01, Created)
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
 DT 01-JUN-2000 (TREMBLrel. 14, Last annotation update)
 DE SIMILARITY TO 9 AMINO ACID REPEATS IN GALACTOSE SPECIFIC LECTINS.
 GN F1865.2.
 OS Caenorhabditis elegans.
 OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditidae;
 OC Rhabditidae; Felodertinae; Caenorhabditis.
 OX NCBI_TaxID=6239;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=94150718; PubMed=7906398;
 RA Wilson R., Ainscough R., Anderson K., Baynes C., Berts M.,
 RA Bonfield J., Burton J., Connell M., Copsey T., Cooper J., Coulson A.,
 RA Craxton M., Dear S., Du Z., Durbin R., Favello A., Fulton L.,
 RA Gardner A., Green P., Hawkins T., Hillier L., Jier M., Johnston L.,
 RA Jones M., Kershaw J., Kirsten J., Laister N., Latelle P.,
 RA Lightning J., Lloyd C., McMurray A., Mortimore B., O'Callaghan M.,
 RA Parsons J., Percy C., Rifkin L., Roopra A., Saunders D., Showkeen R.,
 RA Smaldon N., Smith A., Sonhammer E., Staden R., Sulston J.,
 RA Thierry-Mieg J., Thomas K., Vaudin M., Vaughan K., Waterston R.,
 RA Watson A., Weinstock L., Wilkinson-Sproat J., Wohlman P.,
 RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
 RT elegans."
 RL Nature 366:32-38(1994).
 [2]

RP SEQUENCE FROM N.A.
RA Favellio T.;
RL Submitted (NOV-1995) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RA Waterston R.;
RL Submitted (NOV-1995) to the EMBL/GenBank/DBJ databases.
DR EMBL: U39855; AAA81082.1; -
KW lectin.
SQ SEQUENCE 467 AA; 63169 MW; 7D9BBAB61830431B CRC64;

Query Match 53.8%; Score 49; DB 5; Length 467;
Best Local Similarity 83.3%; Pred. No. 31;
Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 5 WPMWPMW 10
Db 201 WPMWPMW 206

RESULT 14
O26589 PRELIMINARY; PRT; 528 AA.
AC Q26589;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
DE MYOSIN II HEAVY CHAIN (FRAGMENT).
OS Schistosoma mansoni (Blood fluke).
OC Eukaryota; Metazoa; Platyhelminthes; Rhabditophora; Neodermata;
OC Trematoda; Digenea; Strigeidida; Schistosomatidae; Schistosomatidae;
OC Schistosoma.
OX NCBI_TaxID=6183;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=PUERTO RICAN;
RA Amory L.M.;
RL Submitted (APR-1992) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=PUERTO RICAN;
RX MEDLINE=93056536; PubMed=1431131;
RA Scisson L.M., Masterston C.P., Tom T.D., McNally M.T., Lowell G.H.,
RA Strand M.;
RT "Induction of protective immunity in mice using a 62-kDa recombinant
fragment of a Schistosoma mansoni surface antigen.";
RL J. Immunol. 149:3612-3620(1992).
DR EMBL: X65591; CAA46548.1; -
DR HSSP: P08799; 1MMD.
DR InterPro: IPR001637; GlnA_adenyltn.
DR InterPro: IPR000048; IQ.
DR InterPro: IPR001609; myosin_head.
DR InterPro: IPR002928; Myosin_tail.
DR InterPro: IPR000533; Tropomyosin.
DR Pfam: PF00612; IQ; 1.
DR Pfam: PF00663; myosin_head; 1.
DR PRINTS: PR01576; Myosin_tail; 1.
DR PRINTS: PR00194; TROPOMYOSIN.
DR PRODOM: PD000355; myosin_head; 1.
DR SMART: SM00015; IQ; 1.
DR PROSITE: PS00182; GlnA_ADENYLATION; UNKNOWN_1.
DR PROSITE: PS50096; IQ; 1.
KW myosin.
FT NON_TER 1
SQ SEQUENCE 528 AA; 61622 MW; AF075D13EB249B4C CRC64;

Query Match 53.8%; Score 49; DB 5; Length 528;
Best Local Similarity 62.5%; Pred. No. 35;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
OY 1 ILKRWPMW 8

Db 106 VLKRWPMW 113

RESULT 15
O9TY57 PRELIMINARY; PRT; 528 AA.
AC O9TY57;
DT 01-MAY-2000 (TREMBLrel. 13, Created)
DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
DE MYOSIN HEAVY CHAIN (FRAGMENT).
OS Schistosoma japonicum (Blood fluke).
OC Eukaryota; Metazoa; Platyhelminthes; Rhabditophora; Neodermata;
OC Trematoda; Digenea; Strigeidida; Schistosomatidae; Schistosomatidae;
OC Schistosoma.
OX NCBI_TaxID=6182;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CHINESE MAINLAND STRAIN;
RX MEDLINE=99144454; PubMed=990643;
RA Zhang Y.B., Taylor M.G., Bickle Q.D.;
RT "Schistosoma japonicum myosin: cloning, expression and vaccination
studies with the homologue of the S.mansoni myosin fragment IV-5.";
RL Parasite Immunol. 20:583-594(1998).
DR EMBL: U55313; AAC82221.1; -
DR HSSP: P08799; 1MMD.
DR InterPro: IPR000048; IQ.
DR InterPro: IPR001609; myosin_head.
DR InterPro: IPR002928; Myosin_tail.
DR InterPro: IPR000533; Tropomyosin.
DR Pfam: PF00612; IQ; 1.
DR Pfam: PF00663; myosin_head; 1.
DR PRINTS: PR01576; Myosin_tail; 1.
DR PRINTS: PR00194; TROPOMYOSIN.
DR PRODOM: PD000355; myosin_head; 1.
DR SMART: SM00015; IQ; 1.
FT NON_TER 1
SQ SEQUENCE 528 AA; 61406 MW; C54A31F540F5EE05 CRC64;

Query Match 53.8%; Score 49; DB 5; Length 528;
Best Local Similarity 62.5%; Pred. No. 35;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
OY 1 ILKRWPMW 8
Db 106 VLKRWPMW 113

Search completed: January 4, 2002, 08:47:24
Job time: 412 sec

THIS PAGE BLANK (USPTO)

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:47:47 ; Search time 18.1 Seconds
(without alignments)
26.334 Million cell updates/sec

Title: US-09-444-281-35
Perfect score: 91
Sequence: 1 ILKKMPWMPWRRK 13

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 100059 seqs, 36664827 residues
Total number of hits satisfying chosen parameters: 100059

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SwissProt_39.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	73	80.2	144	1 INDC_BOVIN	P33046 bos taurus
2	54	59.3	1173	1 VGL2_CVH22	P15423 human coron
3	49	53.8	492	1 ADRO_BOVIN	P08165 bos taurus
4	49	53.8	715	1 YD55_MYCTU	Q11025 mycobacteri
5	47	51.6	253	1 Y945_MYCTU	P71564 mycobacteri
6	46.5	51.1	1154	1 VGL2_IBVD2	P12222 avian infec
7	46.5	51.1	1162	1 VGL2_IBVB	P11223 avian infec
8	46.5	51.1	1162	1 VGL2_IBVK	P12650 avian infec
9	46.5	51.1	1162	1 VGL2_IBVM	P12651 avian infec
10	46.5	51.1	1163	1 VGL2_IBV6	P05135 avian infec
11	46	50.5	196	1 YA05_SCHPO	Q09677 schizosacch
12	45	49.5	397	1 MMU6_MYCTU	Q10773 mycobacteri
13	45	49.5	505	1 TRE2_PSESS	P21689 pseudomonas
14	45	49.5	512	1 FEN2_YEAST	P25821 saccharomyc
15	45	49.5	964	1 MMU5_MYCTU	O53784 mycobacteri
16	45	49.5	967	1 MMU4_MYCTU	O53735 mycobacteri
17	45	49.5	968	1 MMU2_MYCTU	Q11171 mycobacteri
18	45	49.5	1108	1 CN3B_RAT	O63085 rattus norv
19	45	49.5	1225	1 VGL2_CVPR8	P27655 porcine res
20	45	49.5	1225	1 VGL2_CVPRM	P24413 porcine res
21	45	49.5	1235	1 VGL2_CVPMH	P12423 murine coro
22	45	49.5	1334	1 VGL2_CVMA5	P11224 murine coro
23	45	49.5	1353	1 VGL2_CVHOC	P36334 human coron
24	45	49.5	1363	1 VGL2_CVMBF	P25190 bovine coro
25	45	49.5	1363	1 VGL2_CVBL5	P25191 bovine coro
26	45	49.5	1363	1 VGL2_CVBL6	P25192 bovine coro
27	45	49.5	1363	1 VGL2_CVBM	P15777 bovine coro
28	45	49.5	1363	1 VGL2_CVBO	P25193 bovine coro
29	45	49.5	1363	1 VGL2_CVBV	P25194 bovine coro
30	45	49.5	1376	1 VGL2_CVMA	P22432 murine coro
31	45	49.5	1376	1 VGL2_CVMC	O02385 murine coro
32	45	49.5	1447	1 VGL2_CVPR	Q02167 porcine tira
33	45	49.5	1447	1 VGL2_CVPPU	P07946 porcine tira

34	45	49.5	1447	1 VGL2_CVPR	Q01977 porcine tira
35	45	49.5	1449	1 VGL2_CVPR5	P18450 porcine tira
36	45	49.5	1449	1 VGL2_CVPMI	P33470 porcine tira
37	45	49.5	1451	1 VGL2_CVCAI	P36300 canine ente
38	45	49.5	1452	1 VGL2_FIPV	P10033 feline inte
39	45	49.5	2116	1 MYS2_DICDI	P08799 dictyostell
40	44	48.4	53	1 YDH3_PLAFS	P14589 plasmodium
41	44	48.4	151	1 YBBJ_ECOLI	P75709 escherichia
42	44	48.4	361	1 FUR3_HUMAN	P21217 homo sapien
43	44	48.4	372	1 FUR3_PANTR	O19058 pan troglod
44	44	48.4	451	1 MEN6_ECOLI	P37353 escherichia
45	44	48.4	535	1 YDW6_SCHPO	O13912 schizosacch

ALIGNMENTS

RESULT	1	STANDARD:	PRT:	144	AA.
ID	INDC_BOVIN				
AC	P33046:				
DT	01-OCT-1993 (Rel. 27, Created)				
DT	01-OCT-1993 (Rel. 27, Last sequence update)				
DT	01-NOV-1997 (Rel. 35, Last annotation update)				
DE	INDOLICIDIN PRECURSOR.				
OS	Bos taurus (Bovine).				
OC	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;				
OC	Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;				
OC	Bovidae; Bovinae; Bos.				
OX	NCBI_TaxID=9913;				
RN	[1]				
RP	SEQUENCE FROM N.A.				
RC	TISSUE=Bone marrow:				
RX	MEDLINE=92392368; PubMed=1520337;				
RA	del Sal G., Storici P., Schneider C., Romeo D., Zanetti M.;				
RT	"CDNA cloning of the neutrophil bactericidal peptide indolicidin.";				
RL	Biochem. Biophys. Res. Commun. 187:467-472(1992).				
RN	[2]				
RP	SEQUENCE OF 131-143.				
RC	TISSUE=Neutrophils;				
RX	MEDLINE=92165771; PubMed=1537821;				
RA	Selsted M.E., Novotny M.J., Morris W.L., Tang Y.-Q., Smith W.;				
RT	Cullor J.S.;				
RL	"Indolicidin, a novel bactericidal tridecapeptide amide from neutrophils.";				
CC	J. Biol. Chem. 267:4292-4295(1992).				
CC	- FUNCTION: POTENT MICROBICIDAL ACTIVITY, ACTIVE AGAINST STAPHYLOCOCCUS AUREUS AND ESCHERICHIA COLI.				
CC	- TISSUE SPECIFICITY: LARGE GRANULES OF NEUTROPHILS.				
CC	- PTM: ELASTASE MIGHT BE RESPONSIBLE FOR ITS MATURATION.				
CC	- SIMILARITY: BELONGS TO THE CATHELICIDIN FAMILY.				
CC	-----				
CC	This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See http://www.isb-sib.ch/announce/ or send an email to license@isb-sid.ch).				
CC	-----				
CC	EMBL: X67340; CAA47755.1; -				
DR	PIR: JCI222; JCI222.				
DR	PIR: A42387; A42387.				
DR	InterPro: IPR001894; Cathelicidin.				
DR	Pham: PF00666; Cathelicidins; 1.				
DR	ProDom: PD001838; Cathelicidins; 1.				
DR	PROSITE: PS00946; CATHELICIDINS_1; 1.				
DR	PROSITE: PS00947; CATHELICIDINS_2; 1.				
KW	Antibiotic; Amidation; Signal.				
FT	SIGNAL 1 29				POTENTIAL.
FT	PROPEP 30 130				
FT	PEPTIDE 131 143				INDOLICIDIN.
FT	MOD_RES 30 30				PYRROLIDONE CARBOXYLIC ACID (BY

FT		SIMILARITY).
FT	85	BY SIMILARITY.
FT	107	BY SIMILARITY.
FT	143	AMIDATION (G-144 PROVIDE AMIDE GROUP)
50	144 AA; 16479 MM; E3B1CB8E55C09911 CRC64;	

Query Match	80.28;	Score 73;	DB 1;	Length 144;
Best Local Similarity	100.0%;	Pred. NO. 0.00094;		
Matches	9;	Conservative	0;	Mismatches 0;
			Indels	0;
			Gaps	0;

QY	4	KWPWPWPWRR	12
Db	135	KWPWPWPWRR	143

RESULT	2	
VGL2_CVH22		
ID	VGL2_CVH22	STANDARD;
		PRT; 1173 AA

DT	01-APR-1990 (Rel. 14, Created)
DT	01-APR-1990 (Rel. 14, Last sequence update)
DT	15-JUL-1999 (Rel. 38, Last annotation update)
DE	ES GLYCOPROTEIN PRECURSOR (SPIKE GLYCOPROTEIN) (PEPDOMER PROTEIN)
GN	S.
OS	Human coronavirus (strain 229E).
OC	Viruses; ssRNA positive strand viruses, no DNA stage, Nidovirales
CC	Coronaviridae; Coronavirus.
NCBI_TaxID=11137;	

RP SEQUENCE FROM N.A., PubMed:2345367,
RX MEDLINE-90264837,
RA Raabe T., Schelle-prinz B., Siddell S.G.,
RT Nucleotide sequence of the gene encoding the spike glycoprotein of
RL human coronavirus HCoV 229E.
RT J. Gen. Virol. 71:1065-1073(1990).
CC -I- FUNCTION: THE PERIOMER PROTEIN MEDIATES THE BINDING OF VIRIONS
CC TO THE HOST CELL RECEPTOR AND IS INVOLVED IN MEMBRANE FUSION
CC AND IN SYNCYTIIUM FORMATION.
CC -I- SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN.

CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation
CC the European Bioinformatics Institute. There are no restrictions on lists
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).

Accession	Protein Name	Length (aa)	Source	Function
DR EMBL: X15816; CA34723.1; -	EMBL: X15816; CA34723.1; -			
DR PIR: A34766; VGIHHC.	PIR: A34766; VGIHHC.			
DR InterPro: IPR002551; Corona_S1.	InterPro: IPR002551; Corona_S1.			
DR InterPro: IPR002552; Corona_S2.	InterPro: IPR002552; Corona_S2.			
DR Pfam: PF01600; Corona_S1; 1.	Pfam: PF01600; Corona_S1; 1.			
DR Pfam: PF01601; Corona_S2; 1.	Pfam: PF01601; Corona_S2; 1.			
KM Glycoprotein; Envelope protein; Transmembrane; Signal.	Glycoprotein; Envelope protein; Transmembrane; Signal.			
FT Signal	Signal	1		
FT CHAIN	CHAIN	16		
FT DOMAIN	DOMAIN	16		
FT TRANSMEM	TRANSMEM	1116		
FT DOMAIN	DOMAIN	1136		
FT DOMAIN	DOMAIN	1136		
FT CARBOHYD	CARBOHYD	1136		
FT CARBOHYD	CARBOHYD	23		
FT CARBOHYD	CARBOHYD	62		
FT CARBOHYD	CARBOHYD	98		
FT CARBOHYD	CARBOHYD	147		
FT CARBOHYD	CARBOHYD	171		
FT CARBOHYD	CARBOHYD	176		
FT CARBOHYD	CARBOHYD	220		
FT CARBOHYD	CARBOHYD	243		
FT CARBOHYD	CARBOHYD	326		
FT CARBOHYD	CARBOHYD	333		
FT CARBOHYD	CARBOHYD	440		

FT	CARBOHD	464	464	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	518	518	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	538	538	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	542	542	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	568	568	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	581	581	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	587	587	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	663	663	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	671	671	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	930	930	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	1015	1015	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	1020	1020	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	1037	1037	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	1049	1049	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	1061	1061	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	1066	1066	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	1076	1076	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	1082	1082	N-LINKED (GLCNAC. .)	(POTENTIAL)
FT	CARBOHYD	1096	1096	N-LINKED (GLCNAC. .)	(POTENTIAL)
5Q	SEQUENCE	1173	128639	AA: B9CA2A1A796B3BD	CRC64;

Query Match	59.3%	Score 54;	DB 1;	Length 1173;
Best Local Similarity	85.7%	Pred. No. 2.5;		
Matches	6;	Conservative	0;	Mismatches 1;
			Indels 0;	Gaps 0;

QY	4	KWPWPW	10
Db	1113	KWPWPW	1115

RESULT	3	
ADRO_BOVIN		
ID	ADRO_BOVIN	STANDARD:
AC	P08165:	PRT: 492 AA.
DT	01-AUG-1988 (Rel. 08, Created)	
DT	15-JUL-1998 (Rel. 36, Last sequence update)	
DT	20-AUG-2001 (Rel. 40, Last annotation update)	
DE	NADPH:ADRENODOXIN OXIDOREDUCTASE, MITOCHONDRIAL PRECURSOR (EC 1.18.1.2) (ARENODOXIN REDUCTASE) (AR) (FERREDOXIN-NADP(+	

GN EDXR OR ADXR.
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Cranialta; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla, Ruminantia, Pecora; Bovidae
OC Bovidae; Bovinae, Bos.
OX
NCBI_taxid=9913;

RP SEQUENCE FROM N.A., AND ALTERNATIVE SPLICING.
RA MEDLINE=94177140; PubMed=8130767;
RX Taketa Y., Sagara Y., Kono A., Sekimizu K., Horiuchi T.,
RT "Gene structure of bovine adrenodoxin reductase.";
RL Biol. Pharm. Bull. 16:1200-1206(1993).
RN [2]
RP SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.
RX MEDLINE=88198050; PubMed=3448086;
RA Sagara Y., Taketa Y., Miyata T., Hara T., Horiuchi T.,
RT "Cloning and sequence analysis of adrenodoxin reductase cDNA from
RL bovine adrenal cortex.";
RL J. Biochem 102:1333-1336(1987).

RP SEQUENCE FROM N.A.
RX MEDLINE=87270696; PubMed=303809;
RA Nonaka Y., Murakami H., Yabusaki Y., Kuramitsu S., Kagamiyama H.,
RA Yamano T., Okamoto M.;
RT "Molecular cloning and sequence analysis of full-length cDNA for mRNA
of adrenodoxin oxidoreductase from bovine adrenal cortex.";
RT Blochm. Biophys. Res. Commun. 145:1239-1247(1987).

RP SEQUENCE FROM N.A.
RC TISSUE=Adrenal cortex;
RX MEDLINE=89170752; PubMed=2924777;
RA Hanukoglu I., Gutfinger T.;

RT "cDNA sequence of adrenodoxin reductase. Identification of NADP-binding sites in oxidoreductases.";
 RL Eur. J. Biochem. 180:479-484(1989).
 RN [5]
 RP SEQUENCE OF N-TERMINUS, AND PARTIAL SEQUENCE.
 RC TISSUE-Adrenal cortex;
 RX MEDLINE=88082777; PubMed=3691502;
 RA Hanukoglu I., Gutfinger T., Hanin M., Shively J.E.;
 RT "Isolation of a cDNA for adrenodoxin reductase (ferredoxin-NADP+ reductase). Implications for mitochondrial cytochrome P-450 systems.";
 RL Eur. J. Biochem. 169:449-455(1987).
 RN [6]
 RP X-RAY CRYSTALLOGRAPHY (1.7 ANGSTROMS) OF 33-492.
 RC TISSUE-Adrenal gland;
 RX MEDLINE=99299392; PubMed=10369776;
 RA Ziegler G.A., Vornrhein C., Hanukoglu I., Schulz G.E.;
 RT "The structure of adrenodoxin reductase of mitochondrial P450 systems: election transfer for steroid biosynthesis.";
 RL J. Mol. Biol. 289:981-990(1999).
 CC -1- FUNCTION: SERVES AS THE FIRST ELECTRON TRANSFER PROTEIN IN ALL THE MITOCHONDRIAL P450 SYSTEMS, INCLUDING CHOLESTEROL SIDE CHAIN CLEAVAGE IN ALL STEROIDGENIC TISSUES, STEROID 11-BETA HYDROXYLATION IN THE ADRENAL CORTEX, 25-OH-VITAMIN D3-24 HYDROXYLATION IN THE KIDNEY, AND STEROL C-27 HYDROXYLATION IN THE LIVER.
 CC -1- CATALYTIC ACTIVITY: REDUCED ADRENODOXIN + NADP(+) = OXIDIZED ADRENODOXIN + NADPH.
 CC -1- COFACTOR: FAD.
 CC -1- PATHWAY: CHOLESTEROL SIDE-CHAIN-CLEAVAGE SYSTEM.
 CC -1- SUBUNIT: MONOMER.
 CC -1- SUBCELLULAR LOCATION: MITOCHONDRIAL MATRIX.
 CC -1- ALTERNATIVE PRODUCTS: 2 ISOFORMS; A SHORT FORM (SHOWN HERE) AND A LONG FORM; ARE PRODUCED BY ALTERNATIVE SPLICING. THE LONG FORM REPRESENTS 10-20% OF ALL ADRENODOXIN REDUCTASE MRNA. AND SEEMS TO BE INACTIVE.
 CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - CC the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/> or send an email to license@sib-sib.ch).
 CC
 DR EMBL: M17029; AAA0362.1; -;
 DR EMBL: D00211; BAA00150.1; -;
 DR EMBL: X13736; CAA32002.1; -;
 DR PIR: A29604; A29604.
 DR PIR: J50390; J50390.
 DR PIR: S03558; S03558.
 DR PIR: J70751; J70751.
 DR PDB: 1CJC; 12-APR-99.
 DR PDB: 1ELI; 02-JUN-00.
 DR InterPro: IPR000759; Adnrx_reductase.
 DR PRINTS: PR00419; ADXRDPAE.
 KW Electron transport; Oxidoreductase; Flavoprotein; NADP; FAD;
 KW Mitochondrion; Transit peptide; Alternative splicing; 3D-structure.
 FT TRANSIT 1 32
 FT CHAIN 33 492
 FT VARSPLIT 204 204
 FT VARSPLIT 77 77
 FT CONFLICT 81 94
 FT CONFLICT 124 128
 FT CONFLICT 268 268
 FT CONFLICT 317 318
 FT CONFLICT 333 333
 FT CONFLICT 341 352
 FT SEQUENCE 492 AA; 54338 MW; E68F6F5D18F53131 CRC64;
 Query Match 53.8%; Score 49; DB 1; Length 492;
 Best Local Similarity 43.3%; Pred. No. 5.4;

Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 5 WPMPWP 10
 Db 6 WRMPWP 11
 RESULT 4
 ID YD55_MYCTU STANDARD; PRT; 715 AA.
 AC Q11025;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DE 20-AUG-2001 (Rel. 40, Last annotation update)
 DE HYPOPHETICAL 78.2 KDA PROTEIN RV1355C.
 GN RV1355C OR MT1398 OR MYCY02B10.19C.
 OS Mycobacterium tuberculosis.
 OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
 OX NCBI_TaxID=1773;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=H37RV;
 RX MEDLINE=98295987; PubMed=9634230;
 RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D., Gordon S.V., Eigmeier K., Gas S., Barry C.E. III, Tekala F., Badcock K., Basham D., Brown D., Chillingworth T., Connor R., Davies R., Devlin K., Felwell T., Gentles S., Hamlin N., Holroyd S., Hornsby T., Jagels K., Krogh A., McLean A., Moule S., Murphy L., Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J., Rutter S., Seeger K., Skelton S., Squares S., Squares R., Sultun J.E., Taylor K., Whitehead S., Barrell B.G.;
 RA "Deciphering the biology of Mycobacterium tuberculosis from the complete genome sequence.";
 RT Nature 393:537-544(1998).
 RL [2]
 RN SEQUENCE FROM N.A.
 RP STRAIN=CDC 1551 / Oshkosh;
 RC Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O., Peterson J., Deboy R., Dodson R., Gwinn M.L., Haft D., Hickey E., Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L., Ralcheva A., Ustebach T., Weidman J., Khouri H., Gill J., Mikula A., Bishai W.;
 RT "Whole genome comparison of Mycobacterium tuberculosis clinical and laboratory strains.";
 RL Submitted (Apr-2001) to the EMBL/Genbank/DBSJ databases.
 CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - CC the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/> or send an email to license@sib-sib.ch).
 CC
 DR EMBL: Z75555; CAA99988.1; -;
 DR EMBL: AE007012; AAK45661.1; ALT_INIT.
 DR TIGR: MT1398;
 DR TubercuList; RV1355C;
 DR InterPro: IPR000594; Thif_family.
 DR Pfam: PF00899; Thif_family; 1.
 KW Hypothetical protein; Complete proteome.
 FT SEQUENCE 715 AA; 78181 MW; 455495248A56041C CRC64;
 Query Match 53.8%; Score 49; DB 1; Length 715;
 Best Local Similarity 60.0%; Pred. No. 7.6;
 Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
 Qy 3 KRWMPWMP 12
 Db 64 KRWMPWMP 73

Y	Y945	MYCTU	5	RESULT
AC	Y945	MYCTU	STANDARD:	PRT; 253 AA.
AC	P71564;			
DT	01-NOV-1997	(Rel. 35,	Created)	
DT	01-NOV-1997	(Rel. 35,	Last sequence update)	
DT	20-AUG-2001	(Rel. 40,	Last annotation update)	
DE	PURATIVE OXIDOREDUCTASE RV0945	(EC 1.-.-.-)		
GN	RV0945 OR MT0971 OR MTCY10D7.29C.			
OS	Mycobacterium tuberculosis.			
OC	Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;			
OC	Actinomycetales; Corynebacteriineae; Mycobacteriaceae; Mycobacterium.			
OX	NCBI_TaxId=1773;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	STRAIN-H37RV:			
RX	MEDLINE:98295987; PubMed=9634230;			
RA	Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,			
RA	Gordon S.V., Eiglmeier K., Gas S., Barry C.E. III, Tekala F.,			
RA	Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,			
RA	Davies R., Devlin K., Felwell T., Gentles S., Hamlin N., Holroyd S.,			
RA	Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,			
RA	Oliver S., Seeger K., Skelton S., Squares S., Squares R.,			
RA	Rutter S., Seeger K., Skelton S., Squares S., Squares R.,			
RA	Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;			
RT	"Deciphering the biology of Mycobacterium tuberculosis from the			
RT	complete genome sequence."			
RL	Nature 393:537-544(1998).			
RN	[2]			
RP	SEQUENCE FROM N.A.			
RC	STRAIN-CDC 1551 / Oshkosh;			
RA	Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,			
RA	Petersen J., Deboy R., Dodson R., Gwinn M.L., Haft D., Hickey E.,			
RA	Kolonyak J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L.,			
RA	Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A.,			
RA	Bisbal W.;			
RT	"Whole genome comparison of Mycobacterium tuberculosis clinical and			
RT	laboratory strains."			
RL	Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.			
CC	-I SIMILARITY: BELONGS TO THE SHORT-CHAIN DEHYDROGENASES/REDUCTASES			
CC	(SBR) FAMILY.			
CC	-----			
CC	This SWISS-PROT entry is copyright. It is produced through a collaboration			
CC	between the Swiss Institute of Bioinformatics and the EMBL outstation -			
CC	the European Bioinformatics Institute. There are no restrictions on its			
CC	use by non-profit institutions as long as its content is in no way			
CC	modified and this statement is not removed. Usage by and for commercial			
CC	entities requires a license agreement (see http://www.isb-sib.ch/announce/			
CC	or send an email to license@isb-sib.ch).			
CC	-----			
RR	EMBL; Z79700; CAB02005.1; -			
RR	EMBL; AE006982; AAK45219.1; -			
DR	TIGR; MT0971; -			
DR	TubercuList; RV0945; -			
DR	InterPro: IPR002198; ADH_short.			
DR	Pfam; PF00106; adh_short; 1.			
DR	PROSITE; PS00061; ADH_SHORT; 1.			
KW	Hypothetical protein; Oxidoreductase; Complete proteome.			
FT	ACT_SITE 159 159			
FT	BY SIMILARITY			
SO	SEQUENCE 253 AA; 27138 MW; BAD937208842DA12 CRC64;			
Y	Query Match			
Y	Best Local Similarity 51.6%; Score 47; DB 1; Length 253;			
Y	Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;			
Y	6 PWM PWM 10			
Y				
Y	230 PWM PWM 234			

[illegible]

SQ SEQUENCE 1154 AA; 127502 MW; D79F37AF89F1A37F CRC64;
 Query Match 51.1%; Score 46.5; DB 1; Length 1154;
 Best Local Similarity 61.5%; Pred. No. 26;
 Matches 8; Conservative 1; Mismatches 1; Indels 3; Gaps 1;

OY 1 ILK---KMPMPW 10
 ||| |||||
 Db 1086 ILKTYIKMPWYVW 1098

RESULT 7
 VGL2_IBVK STANDARD; PRT; 1162 AA.
 ID P11223; P05134;
 AC 01-JUL-1989 (Rel. 11, Created)
 DT 01-JUL-1989 (Rel. 11, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE E2 GLYCOPROTEIN PRECURSOR (SPIKE GLYCOPROTEIN) (PEPLOMER PROTEIN)
 DE [CONTAINS: SPIKE PROTEIN S1; SPIKE PROTEIN S2].
 GN S.
 OS Avian infectious bronchitis virus (strain Beaudette) (IBV).
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
 OC Coronaviridae; Coronavirus.
 OX NCBI_TaxID=11122;
 RX MEDLINE FROM N.A. PubMed=2984314;
 RA Binn M.M., Boursnell M.E.G., Cavanagh D., Pappind D.J.C.,
 RA Brown T.D.K.;
 RT "Cloning and sequencing of the gene encoding the spike protein of the
 RT coronavirus IBV";
 RL J. Gen. Virol. 66:719-726(1985).
 RN [2]
 RP SEQUENCE FROM N.A. PubMed=3025348;
 RX MEDLINE=87085499; PubMed=3025348;
 RA Binn M.M., Boursnell M.E.G., Tomley F.M., Brown D.K.;
 RT "Comparison of the spike precursor sequences of coronavirus IBV
 RT strains M41 and 6/82 with that of IBV Beaudette";
 RL J. Gen. Virol. 67:2825-2831(1986).
 CC -1- FUNCTION: THE PELOMER PROTEIN MEDIATES THE BINDING OF VIRIONS
 CC TO THE HOST CELL RECEPTOR AND IS INVOLVED IN MEMBRANE FUSION.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL; M95169; AAA70235.1; -;
 DR EMBL; X02342; CAA26201.1; -;
 DR InterPro: IPR002551; Corona_S1.
 DR InterPro: IPR002552; Corona_S2.
 DR Pfam: PF01600; Corona_S1; 1.
 DR Pfam: PF01601; Corona_S2; 1.
 KW Glycoprotein; Envelope protein; Transmembrane; Signal.
 FT SIGNAL 1 18
 FT CHAIN 19 1162 E2 GLYCOPROTEIN.
 FT CHAIN 19 537 SPIKE PROTEIN S1.
 FT DOMAIN 1120 1137 SPIKE PROTEIN S2.
 FT CARBOHYD 51 91 CYS-RICH.
 FT CARBOHYD 77 79 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 103 103 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 144 144 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 163 163 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 178 178 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 212 212 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 237 237 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 247 247 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 264 264 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 276 276 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 306 306 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 425 425 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 447 447 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 513 513 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 530 530 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 579 579 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 591 591 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 669 669 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 714 714 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 947 947 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 960 960 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 979 979 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1014 1014 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1038 1038 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1051 1051 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1074 1074 N-LINKED (GLCNAC. . .) (POTENTIAL).
 SQ SEQUENCE 1162 AA; 128046 MW; 0BAAD58113C8EBD5 CRC64;

OY 1 ILK---KMPMPW 10
 ||| |||||
 Db 1085 ILKTYIKMPWYVW 1097

Query Match 51.1%; Score 46.5; DB 1; Length 1162;
 Best Local Similarity 61.5%; Pred. No. 26;
 Matches 8; Conservative 1; Mismatches 1; Indels 3; Gaps 1;

RESULT 8
 VGL2_IBVK STANDARD; PRT; 1162 AA.
 ID P12650;
 AC 01-OCT-1989 (Rel. 12, Created)
 DT 01-OCT-1989 (Rel. 12, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE E2 GLYCOPROTEIN PRECURSOR (SPIKE GLYCOPROTEIN) (PEPLOMER PROTEIN)
 DE [CONTAINS: SPIKE PROTEIN S1; SPIKE PROTEIN S2].
 GN S.
 OS Avian infectious bronchitis virus (strain KB8523) (IBV).
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
 OC Coronaviridae; Coronavirus.
 OX NCBI_TaxID=11126;
 RX MEDLINE FROM N.A. PubMed=2841803;
 RA Suto S., Sato S., Okabe T., Nakai M., Sasaki N.;
 RT "Cloning and sequencing of genes encoding structural proteins of
 RT avian infectious bronchitis virus";
 RL Virology 165:589-595(1988).
 CC -1- FUNCTION: THE PELOMER PROTEIN MEDIATES THE BINDING OF VIRIONS
 CC TO THE HOST CELL RECEPTOR AND IS INVOLVED IN MEMBRANE FUSION.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL; M21515; AAA66578.1; -;
 DR PIR; B29249; VG1HAK.
 DR InterPro: IPR002551; Corona_S1.
 DR InterPro: IPR002552; Corona_S2.
 DR Pfam: PF01600; Corona_S1; 1.
 DR Pfam: PF01601; Corona_S2; 1.
 KW Glycoprotein; Envelope protein; Transmembrane; Signal.
 FT SIGNAL 1 18
 FT CHAIN 19 1162 E2 GLYCOPROTEIN.
 FT CHAIN 19 537 SPIKE PROTEIN S1.

```

FT CHAIN 538 1162 SPIKE PROTEIN S2.
FT DOMAIN 1120 1137 CYS-RICH.
FT CARBOHYD 51 81 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 77 77 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 103 103 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 144 144 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 163 163 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 178 178 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 212 212 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 237 237 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 247 247 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 264 264 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 271 271 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 276 276 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 306 306 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 425 425 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 447 447 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 513 513 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 530 530 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 579 579 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 591 591 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 669 669 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 676 676 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 714 714 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 947 947 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 960 960 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 979 979 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1014 1014 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1051 1051 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1058 1058 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1074 1074 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 1162 AA; 128537 MW; 22930363597EAB8F CRC64;

```

Query Match .51.1%; Score 46.5; DB 1; Length 1162;

Best Local Similarity 61.5%; Pred. No. 26; Mismatches 1; Indels 3; Gaps 1;

```

Oy 1 ILK---KWPMPW 10
    ||| ||||: |
Db 1085 ILKTYIKWPWYV 1097

```

RESULT 9

VG12_IBVM STANDARD; PRT; 1162 AA.

AC P12651; 01-OCT-1989 (Rel. 12, Created)

DT 01-OCT-1989 (Rel. 12, Last sequence update)

DE E2 GLYCOPROTEIN PRECURSOR (SPIKE GLYCOPROTEIN) (PEPLOMER PROTEIN)

[CONTAINS: SPIKE PROTEIN S1; SPIKE PROTEIN S2].

GN S.

OS Avian infectious bronchitis virus (strain M41) (IBV).

OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;

OC Coronaviridae; Coronavirus.

OX NCBI_Taxid=11127;

RN [1]

RP MEDLINE=87021475; PubMed=2429473;

RX Nieuwers H.G.M., Ienstra J.A., Spaan W.J.M., Zijderfeld A.J.,

Blumink-Plum N.M.C., Hong F., van Scharrenburg G.J.M.,

RA Horzinek M.C., van der Zeijst B.A.M.;

RT "The peplomer protein sequence of the M41 strain of coronavirus IBV

and its comparison with Beaudette strains.";

RL Virus Res. 5:253-263(1986).

CC -1- FUNCTION: THE PELOMER PROTEIN MEDIATES THE BINDING OF VIRIONS

CC TO THE HOST CELL RECEPTOR AND IS INVOLVED IN MEMBRANE FUSION.

CC This SWISS-PROT entry is copyright. It is produced through a collaboration

CC between the Swiss Institute of Bioinformatics and the EMBL Outstation

CC the European Bioinformatics Institute. There are no restrictions on its

CC use by non-profit institutions as long as its content is in no way

CC modified and this statement is not removed. Usage by and for commercial

CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>

CC or send an email to license@isb-sib.ch).

CC -----

DR EMBL; M21883; AAA65575.1; -;

DR EMBL; A24863; CA01736.1; -;

DR PIR; S07421; S07421.

DR InterPro; IPR002551; Corona_S1.

DR InterPro; IPR002552; Corona_S2.

DR Pfam; PF01600; Corona_S1; 1.

DR Pfam; PF01601; Corona_S2; 1.

KW Glycoprotein; Envelope protein; Transmembrane; Signal.

FT SIGNAL 1 18

FT CHAIN 19 1162 E2 GLYCOPROTEIN S1.

FT CHAIN 538 1162 SPIKE PROTEIN S2.

FT CHAIN 1120 1137 CYS-RICH.

FT DOMAIN 51 51 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 77 77 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 103 103 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 144 144 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 163 163 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 178 178 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 212 212 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 237 237 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 247 247 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 264 264 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 271 271 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 276 276 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 306 306 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 425 425 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 447 447 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 513 513 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 530 530 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 579 579 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 591 591 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 669 669 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 714 714 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 947 947 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 960 960 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 979 979 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 1014 1014 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 1038 1038 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 1051 1051 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 1074 1074 N-LINKED (GLCNAC. . .) (POTENTIAL).

SQ SEQUENCE 1162 AA; 128077 MW; 3C9CC70938492DDA CRC64;

Query Match .51.1%; Score 46.5; DB 1; Length 1162;

Best Local Similarity 61.5%; Pred. No. 26; Mismatches 1; Indels 3; Gaps 1;

```

Oy 1 ILK---KWPMPW 10
    ||| ||||: |
Db 1085 ILKTYIKWPWYV 1097

```

RESULT 10

VG12_IBV6 STANDARD; PRT; 1163 AA.

AC P05135; 13-AUG-1987 (Rel. 05, Created)

DT 13-AUG-1987 (Rel. 05, Last sequence update)

DT 20-AUG-2001 (Rel. 40, Last annotation update)

DE E2 GLYCOPROTEIN PRECURSOR (SPIKE GLYCOPROTEIN) (PEPLOMER PROTEIN)

[CONTAINS: SPIKE PROTEIN S1; SPIKE PROTEIN S2].

GN S.

OS Avian infectious bronchitis virus (strain 6/82) (IBV).

OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;

OC Coronaviridae; Coronavirus.

OX NCBI_Taxid=11121;

RN [1]

RP SEQUENCE FROM N.A.
RX MEDLINE-87085499; PubMed-3025348;
RA Bins M.M., Bournell M.E.G., Tomley F.M., Brown D.K.;
RT "Comparison of the spike precursor sequences of coronavirus IBV
RL strains M41 and 6/82 with that of IBV Beaudette.";
CC J. Gen. Virol. 67:2825-2831(1986).
CC -I- FUNCTION: THE PEPLIMER PROTEIN MEDIATES THE BINDING OF VIRIONS
CC TO THE HOST CELL RECEPTOR AND IS INVOLVED IN MEMBRANE FUSION.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; X04723; CAA28432.1; -
DR InterPro: IPR002551; Corona_S1.
DR InterPro: IPR002552; Corona_S2.
DR Pfam: PF01600; Corona_S1; 1.
DR Pfam: PF01601; Corona_S2; 1.
KW Glycoprotein; Transmembrane; Signal.
FT SIGNAL 1 18
FT CHAIN 19 1163 E2 GLYCOPROTEIN.
FT CHAIN 19 538 SPIKE PROTEIN S1.
FT CHAIN 539 1163 SPIKE PROTEIN S2.
FT DOMAIN 1121 1138 CYS-RICH.
FT CARBOHYD 23 23 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 51 51 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 74 74 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 102 102 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 139 139 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 145 145 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 164 164 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 179 179 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 213 213 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 238 238 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 248 248 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 265 265 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 272 272 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 277 277 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 307 307 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 426 426 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 448 448 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 514 514 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 531 531 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 543 543 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 580 580 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 592 592 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 670 670 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 677 677 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 948 948 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 961 961 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 980 980 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1015 1015 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1039 1039 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1052 1052 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1075 1075 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 1163 AA; 128684 MW; 8FE344CF2995A78C CRC64;

Query Match 51.1%; Score 46.5; DB 1; Length 1163;
Best Local Similarity 61.5%; Pred. No. 26;
Matches 8; Conservative 1; Mismatches 1; Indels 3; Gaps 1;

OY 1 ILK---KMPMPV 10
DB 1086 ILKTYIKMPYV 1098

RESULT 11
YA05_SCHPO

ID YA05_SCHPO STANDARD; PRT; 196 AA.
AC 009677;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 01-NOV-1995 (Rel. 32, Last annotation update)
DE HYPOTHETICAL 22.1 KDA PROTEIN C5H10.05C IN CHROMOSOME I.
GN SPAC5H10.05C.
OS Schizosaccharomyces pombe (fission yeast).
OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
OC Schizosaccharomycetales; Schizosaccharomycetaceae;
OC Schizosaccharomycetes.
OX NCBI_TaxID=4896;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-972;
RA Connor R., Churcher C.M., Barrell B.G., Rajandream M.A., Walsh S.V.;
RL Submitted (MAY-1995) to the EMBL/Genbank/DBJ databases.
CC -I- SIMILARITY: STRONG, TO BACTERIAL MODULATOR OF DRUG ACTIVITY B
CC (MDAB).
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; Z49811; CAA89955.1; -
DR InterPro: IPR003680; NADHdh_2.
DR Pfam: PF02525; NADHdh_2; 1.
KW Hypothetical protein.
SQ SEQUENCE 196 AA; 22104 MW; 436764DA9E26074C CRC64;

Query Match 50.5%; Score 46; DB 1; Length 196;
Best Local Similarity 47.1%; Pred. No. 5.9;
Matches 8; Conservative 3; Mismatches 2; Indels 4; Gaps 2;

OY 1 ILKWP-WW---PMRRK 13
DB 62 ILYWPGWMMGTWKLK 78

RESULT 12
ID MML6_MYCTU STANDARD; PRT; 397 AA.
AC Q10773;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE PUTATIVE MEMBRANE PROTEIN MML6.
GN MML6 OR RV1557 OR MFL608 OR MTCV48.08C.
OS Mycobacterium tuberculosis.
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1773;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-H37RV;
RX MEDLINE-98295987; PubMed-9634230;
RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
RA Gordon S.V., Eigmeier K., Gas S., Barry C.E. III, Tejaia F.,
RA Badoock K., Basham D., Brown D., Chillingworth T., Connor R.,
RA Davies R., Devlin K., Feltwell T., Gentles S., Hamlin N., Holtroyd S.,
RA Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
RA Rutter S., Seeger K., Skellon S., Squares S., Squares R.,
RA Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;
RT "Deciphering the biology of Mycobacterium tuberculosis from the
RT complete genome sequence.";
RL Nature 393:537-544(1998).
RN [2]

```

RP SEQUENCE FROM N.A.
RC STRAIN=CDC 1551 / Oshkosh;
RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
RA Peterson J., Deboy R., Dodson R., Gwin M.L., Haft D., Hickey E.,
RA Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L.,
RA Delcher A., Utterback T., Weidman J., Kouri H., Gill J., Mikula A.,
RA Biswal W.;
RT "Whole genome comparison of Mycobacterium tuberculosis clinical and
RT laboratory strains."
RL Submitted (Apr-2001) to the EMBL/Genbank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
CC -1- SIMILARITY: BELONGS TO THE MPM1 FAMILY.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (see http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; Z74020; CAA98334.1; -
DR EMBL; AE007027; AAK45875.1; -
DR TIGR; MT1608; -
DR TubercuList; RV1557; -
KW Hypothetical protein; Transmembrane; Complete proteome.
FT TRANSMEM 161 181 POTENTIAL.
FT TRANSMEM 190 210 POTENTIAL.
FT TRANSMEM 214 234 POTENTIAL.
FT TRANSMEM 242 262 POTENTIAL.
FT TRANSMEM 293 313 POTENTIAL.
FT TRANSMEM 330 350 POTENTIAL.
SQ SEQUENCE 397 AA; 42421 MW; 678DC86E24472BF4 CRC64;

Query Match 49.5%; Score 45; DB 1; Length 397;
Best Local Similarity 54.5%; Pred. No. 15;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 ILKWPWPWR 11
Db 348 LIGRWFWPQR 358

RESULT 13
TRPE_PSESS STANDARD; PRT; 505 AA.
AC P21689;
DT 01-MAY-1991 (Rel. 18, Last sequence update)
DT 01-MAY-1991 (Rel. 18, Last sequence update)
DT 30-MAY-2000 (Rel. 39, Last annotation update)
DE ANTHRANILATE SYNTHASE COMPONENT I (EC 4.1.3.27).
GN TRPE.
OS Pseudomonas syringae (pv. savastanoi).
OC Bacteria; Proteobacteria; gamma subdivision; Pseudomonadaceae;
OC Pseudomonas.
OX NCBI_TaxID=29438;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=91100331; PubMed=1987141;
RA da Costa E., Silva O., Kosuge T.;
RT "Molecular characterization and expression analysis of the
RT anthranilate synthase gene of Pseudomonas syringae subsp.
RT savastanoi."
RL J. Bacteriol. 173:463-471(1991).
CC -1- CATALYTIC ACTIVITY: CHORISMATE + L-GLUTAMINE = ANTHRANILATE +
CC PYRUVATE + L-GLUTAMATE.
CC -1- PATHWAY: FIRST STEP IN BIOSYNTHESIS OF TRYPTOPHAN.
CC -1- SUBUNIT: TRIMER OF TWO COMPONENTS I AND TWO COMPONENTS II (BY
CC SIMILARITY).
CC -1- MISCELLANEOUS: COMPONENT I CATALYZES THE FORMATION OF ANTHRANILATE
CC USING AMMONIA RATHER THAN GLUTAMINE, WHEREAS COMPONENT II PROVIDES
CC GLUTAMINE AMIDOTRANSFERASE ACTIVITY.

```

```

CC -1- SIMILARITY: BELONGS TO THE ANTHRANILATE SYNTHASE COMPONENT I
CC FAMILY.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (see http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; M55911; AAA26016.1; -
DR PIR; A39128; A39128.
DR InterPro; IPR000350; Chorismate_bind.
DR Pfam; PF00425; chorismate_bind; 1.
DR PRINTS; PR00095; ANTSNTHASE1.
DR Prodom; PD000779; Chorismate_bind; 1.
KW Tryptophan biosynthesis; Lyase.
SQ SEQUENCE 505 AA; 56084 MW; A38E81931331F6B8 CRC64;

Query Match 49.5%; Score 45; DB 1; Length 505;
Best Local Similarity 71.4%; Pred. No. 19;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 7 WWPWRK 13
Db 485 WWPWR 491

RESULT 14
FEN2_YEAST STANDARD; PRT; 512 AA.
ID FEN2_YEAST
AC P25621;
DT 01-MAY-1992 (Rel. 22, Created)
DT 01-MAY-1992 (Rel. 22, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE PROBABLE TRANSPORTER FEN2.
GN FEN2 OR YCR028C OR YCR28C.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomyces.
OX NCBI_TaxID=4932;
RN [1]
RP SEQUENCE FROM N.A.
RA Cederberg H., Hohmann S., Schaaff-gerstenschlaeger I., Huse K.,
RA Zimmermann F.K.; (1992) to the EMBL/Genbank/DBJ databases.
RL Submitted (MAR-1992) to the EMBL/Genbank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=93070619; PubMed=1332309;
RA Carbone M.L.A., Panzeri L., Falconi M.M., Carcano C., Plevani P.,
RA Lucchini G.;
RT "Nucleotide sequence of 9.2 kb left of CRI1 on yeast chromosome III
RT from strain AB872: evidence for a ty insertion and functional
RT analysis of open reading frame YCR28."
RL Yeast 8:805-812(1992).
RN [3]
RP SIMILARITY TO DAL5 FAMILY.
RX MEDLINE=94147996; PubMed=8313894;
RA Koonin E.V., Bork P., Sander C.;
RT "Yeast chromosome III: new gene functions."
RL EMBO J. 13:493-503(1994).
RN [4]
RP CHARACTERIZATION.
RX MEDLINE=96367594; PubMed=8771708;
RA Marcireau C., Joets J., Poussel D., Guilloton M., Karst F.;
RT "FEN2: a gene implicated in the catabolite repression-mediated
RT regulation of ergosterol biosynthesis in yeast."
RL Yeast 12:531-539(1996).
CC -1- FUNCTION: INVOLVED IN THE CATABOLITE REPRESSION-MEDIATED
CC REGULATION OF ERGOSTEROL BIOSYNTHESIS AND IN FENPROPIOMORPH
CC RESISTANCE.

```

CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE ALLANTOATE PERMEASE FAMILY.
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL: X59720; CAA42320.1;
 DR PIR: S19439; S19439.
 DR PIR: S25336; S25336.
 DR SGD: S0000623; FEN2.
 KW Transmembrane; Transport.
 FT TRANSMEM 28 48 POTENTIAL.
 FT TRANSMEM 80 100 POTENTIAL.
 FT TRANSMEM 103 123 POTENTIAL.
 FT TRANSMEM 133 153 POTENTIAL.
 FT TRANSMEM 165 185 POTENTIAL.
 FT TRANSMEM 199 219 POTENTIAL.
 FT TRANSMEM 272 292 POTENTIAL.
 FT TRANSMEM 313 333 POTENTIAL.
 FT TRANSMEM 343 363 POTENTIAL.
 FT TRANSMEM 373 393 POTENTIAL.
 FT TRANSMEM 402 422 POTENTIAL.
 FT TRANSMEM 435 455 POTENTIAL.
 FT TRANSMEM 485 505 POTENTIAL.
 FT CONFLICT 104 104 W -> V (IN REF. 2).
 FT SEQUENCE 512 AA; 58256 MW; 361942E74C62B3B4 CRC64;

Query Match 49.5%; Score 45; DB 1; Length 512;
 Best Local Similarity 62.5%; Pred. No. 20;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 ILKKWMPW 8
 :|:|:|:|
 Db 268 VLKRWMMW 275

RESULT 15
 MML5_MYCTU STANDARD; PRT: 964 AA.
 AC 053784;
 DT 30-MAY-2000 (Rel. 39, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE PUTATIVE MEMBRANE PROTEIN MML5.
 GN MML5 OR RV0676C OR MT0705 OR MT040.04C.
 OS Mycobacterium tuberculosis.
 CC Bacteria; Filicutes; Actinobacteria; Actinobacteridae;
 CC Actinomycetales; Corynebacteriaceae; Mycobacteriaceae; Mycobacterium.
 CC NCBI_TaxID=1773;
 OX 11
 RN SEQUENCE FROM N.A.
 RP STRAIN=H37RV;
 RC MEDLINE=98295987; PubMed=9634230;
 RX Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
 Gordon S.V., Eigmeier K., Gas S., Barry C.E. III, Tekala F.,
 Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
 Davies R., Devlin K., Feltwell T., Gentles S., Hamlin N., Holroyd S.,
 Hornsby T., Jagels K., Kiroch A., McLean J., Moule S., Murphy L.,
 Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
 Ruter S., Seeger K., Skelton S., Squares S., Squares R.,
 Sultson J.E., Taylor K., Whitehead S., Barrett B.G.;
 RT "Deciphering the biology of Mycobacterium tuberculosis from the
 complete genome sequence."
 RL Nature 393:537-544(1998).
 RN 121
 RP SEQUENCE FROM N.A.
 RC STRAIN=CDC 1551 / Oshkosh;
 RX Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,

RA Peterson J., DeBoy R., Dodson R., Gwinn M.L., Haft D., Hickey E.,
 RA Kolonay J.F., Nelson W.C., Umayan L.A., Ermolaeva M.D., Salzberg S.L.,
 RA Delcher A., Utterback T., Weidman J., Kouri H., Gill J., Mikula A.,
 RA Bishai W.;
 RA "Whole genome comparison of Mycobacterium tuberculosis clinical and
 RA laboratory strains."
 RT Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
 CC -1- SIMILARITY: BELONGS TO THE MML5 FAMILY.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL: AL021943; CAA17459.1;
 DR EMBL: AE006964; AAK4930.1;
 DR TIGR: MT0705;
 DR TubercuList; RV0676C;
 DR InterPro; IPR001036; ACR_tran.
 DR PRINTS; PR00702; ACRIFLAVINRP.
 KW Hypothetical protein; Transmembrane; Complete proteome.
 FT TRANSMEM 31 51 POTENTIAL.
 FT TRANSMEM 203 223 POTENTIAL.
 FT TRANSMEM 230 250 POTENTIAL.
 FT TRANSMEM 255 275 POTENTIAL.
 FT TRANSMEM 302 322 POTENTIAL.
 FT TRANSMEM 340 360 POTENTIAL.
 FT TRANSMEM 389 409 POTENTIAL.
 FT TRANSMEM 445 465 POTENTIAL.
 FT TRANSMEM 494 514 POTENTIAL.
 FT TRANSMEM 543 563 POTENTIAL.
 FT TRANSMEM 572 592 POTENTIAL.
 FT TRANSMEM 601 621 POTENTIAL.
 FT TRANSMEM 630 650 POTENTIAL.
 FT TRANSMEM 659 679 POTENTIAL.
 FT TRANSMEM 688 708 POTENTIAL.
 FT TRANSMEM 717 737 POTENTIAL.
 FT TRANSMEM 746 766 POTENTIAL.
 FT TRANSMEM 775 795 POTENTIAL.
 FT TRANSMEM 804 824 POTENTIAL.
 FT TRANSMEM 833 853 POTENTIAL.
 FT TRANSMEM 862 882 POTENTIAL.
 FT TRANSMEM 891 911 POTENTIAL.
 FT TRANSMEM 920 940 POTENTIAL.
 FT CONFLICT 948 948 I -> V (IN REF. 2).
 FT SEQUENCE 964 AA; 104784 MW; B7C945940A1176BD CRC64;

Query Match 49.5%; Score 45; DB 1; Length 964;
 Best Local Similarity 66.7%; Pred. No. 35;
 Matches 6; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 ILKKWMPW 9
 :|:|:|:|
 Db 932 LLGKWFMP 940

Search completed: January 4, 2002, 08:47:48
 Job time: 406 sec

THIS PAGE BLANK (USPTO)

GenCore version 4.5
Copyright (c) 1993 - 2000, Compugen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:41:31 ; Search time 27.18 Seconds
(without alignments)
36,434 Million cell updates/sec

Title: US-09-444-281-35

Perfect score: 91

Sequence: 1 ILKKPMPMPMRK 13

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 219241 seqs, 76174552 residues

Total number of hits satisfying chosen parameters: 219241

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database:

PIR:68:***
1: p1r1:***
2: p1r2:***
3: p1r3:***
4: p1r4:***

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	73	80.2	144	JC1222	indolicidin precursor
2	59.3	1173	1	VGIRHC	E2 glycoprotein pr
3	56.0	289	2	T12505	hypothetical prote
4	49	53.8	298	B72492	hypothetical prote
5	49	53.8	498	JT0751	ferredoxin--NADP+
6	49	53.8	527	S33068	myosin heavy chain
7	49	53.8	715	B70741	probable meey prot
8	49	53.8	1940	A59287	myosin heavy chain
9	48	52.7	111	T29295	hypothetical prote
10	47.5	52.2	114	T36208	hypothetical prote
11	47	51.6	248	S23449	NADH oxidase (H2O2
12	47	51.6	253	G70715	histidinol-phospha
13	46.5	51.1	352	S77354	peptidomeric polypro
14	46.5	51.1	621	S37664	peptidomeric polypro
15	46.5	51.1	630	S37663	peptidomeric polypro
16	46.5	51.1	1154	VGIRHB	E2 glycoprotein pr
17	46.5	51.1	1162	VGIRAK	E2 glycoprotein pr
18	46.5	51.1	1162	S07421	E2 glycoprotein pr
19	46.5	51.1	1162	S14939	E2 glycoprotein pr
20	46.5	51.1	1162	S14940	E2 glycoprotein pr
21	46	50.5	196	S55483	modulator of drug
22	46	50.5	617	T22175	hypothetical prote
23	46	50.5	623	T22177	hypothetical prote
24	46	50.5	1333	S65812	RNA-directed DNA p
25	45	49.5	273	B82646	monofunctional bio
26	45	49.5	276	B83161	probable short-cha
27	45	49.5	397	B70763	probable membrane
28	45	49.5	448	H72376	hypothetical prote
29	45	49.5	505	A39128	anthranilate synth

30	45	49.5	512	2	S19439	probable membrane
31	45	49.5	964	2	E70826	probable membrane
32	45	49.5	967	2	C70831	probable mmp14 pro
33	45	49.5	968	2	F70746	probable mmp12 pro
34	45	49.5	1108	2	A48508	cyclic-nucleotide
35	45	49.5	1225	1	S24284	E2 glycoprotein pr
36	45	49.5	1225	2	A36607	E2 glycoprotein pr
37	45	49.5	1235	1	VGIRHJ	E2 glycoprotein pr
38	45	49.5	1324	1	VGIRH9	E2 glycoprotein pr
39	45	49.5	1353	1	JQ2168	E2 glycoprotein pr
40	45	49.5	1361	2	S29998	surface protein
41	45	49.5	1362	2	A37474	surface glycoprote
42	45	49.5	1363	1	VGIRHM	E2 glycoprotein pr
43	45	49.5	1363	1	VGIRH9	E2 glycoprotein pr
44	45	49.5	1363	1	VGIRH1	E2 glycoprotein pr
45	45	49.5	1363	1	VGIRH9	E2 glycoprotein pr

ALIGNMENTS

```
RESULT 1
JC1222
indolicidin precursor - bovine
N:Alternate names: antimicrobial peptide
C:Species: Bos primigenius taurus (cattle)
C:Date: 10-Sep-1999 #sequence, revision 10-Sep-1999 #text-change 10-Sep-1999
C:Accession: JC1222; A42387; S25664
R:del Sal, G.; Storici, P.; Schneider, C.; Romeo, D.; Zanetti, M.
Biochem. Biophys. Res. Commun. 187, 467-472, 1992
A:Title: cDNA cloning of the neutrophil bactericidal peptide indolicidin.
A:Reference number: JC1222; WUID:92392368
A:Accession: JC1222
A:Molecule type: mRNA
A:Residues: 1-144 <SAL>
A:Cross-references: EMBL:X67340; NID:q462; PIDN:CAA47755.1; PID:q463
A:Experimental source: bone marrow
R:Selsted, M.E.; Novotny, M.J.; Morris, W.L.; Tang, Y.Q.; Smith, W.; Cullor, J.S.
J. Biol. Chem. 267, 4292-4295, 1992
A:Title: Indolicidin, a novel bactericidal tridecapeptide amide from neutrophils.
A:Reference number: A42387; WUID:92165771
A:Accession: A42387
A:Molecule type: protein
A:Residues: 131-143 <SEL>
A:Experimental source: neutrophils
A>Note: sequence extracted from NCBI backbone (NCBI:83840)
C:Superfamily: cathelin; cystatin homology
C:Keywords: amidated carboxyl end
F:1-29/Domain: signal sequence #status predicted <SIG>
F:22-129/Domain: cystatin homology <CYS>
F:30-110/Domain: propeptide #status predicted <PRO>
F:111-143/Product: indolicidin #status experimental <MNT>
F:143/Modified site: amidated carboxyl end (Arg) (amide in mature form from following
```

Query Match 80.2% Score 73; DB 1; Length 144;
Best Local Similarity 100.0% Pred. No. 0.0027;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 KMPMPMR 12
Db 135 KMPMPMR 143

RESULT 2
VGIRHC
E2 glycoprotein precursor - human coronavirus (strain 229E)
N:Alternate names: peplomer glycoprotein; S glycoprotein; spike glycoprotein
C:Species: human coronavirus
A>Note: host Homo sapiens (man)
C:Date: 31-Dec-1991 #sequence, revision 31-Dec-1991 #text-change 16-Jun-2000
C:Accession: A34766; S05460
R:Raabe, T.; Schelle-Prinz, B.; Siddell, S.G.

J. Gen. Virol. 71, 1065-1073, 1990
A:Title: Nucleotide sequence of the gene encoding the spike glycoprotein of human corona
A:Reference number: A34766; MUID:90264837
A:Accession: A34766
A:Molecule type: mRNA
A:Residues: 1-1173 <RA>
A:Cross-references: EMBL:X16816; NID:958926; PIDN:CAA34723.1; PID:958927
A:Experimental source: strain 229E
R:Raabe, T.; Sidel, S.
Nucleic Acids Res. 17, 6387, 1989
A:Title: Nucleotide sequence of the human coronavirus HCV 229E mRNA 4 and mRNA 5 unique
A:Reference number: A34038; MUID:8936667
A:Accession: S03560
A:Status: translation not shown
A:Molecule type: mRNA
A:Residues: 1159-1173 <RA2>
A:Cross-references: EMBL:X15654; NID:958921; PIDN:CAA33680.1; PID:91334827
C:Superfamily: coronavirus E2 glycoprotein
C:Keywords: glycoprotein; transmembrane protein
F:1-15/Domain: signal sequence #status predicted <SIG>
F:16-1173/Product: E2 glycoprotein #status predicted <MAT>
F:1116-1138/Domain: transmembrane #status predicted <TM>
F:23,62,98,147,171,176,220,243,326,333,440,464,518,538,542,568,581,587,663,671,930,1015,

Query Match
Best Local Similarity 59.3%; Score 54; DB 1; Length 1173;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 4 KMPMPM 10
DB 1113 KMPMPM 1119

RESULT 3
T12505
hypothetical protein DKFZp434C192.1 - human (fragment)
C:Species: Homo sapiens (man)
C>Date: 23-Jul-1999 #sequence_revision 23-Jul-1999 #text_change 23-Jul-1999
R:Ansorge, W.; Winkler, U.; Mewes, H.W.; Gassenhuber, J.; Wiemann, S.
submitted to the Protein Sequence Database, June 1999
A:Reference number: 217527
A:Accession: T12505
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-299 <ANS>
A:Cross-references: EMBL:AL096753
A:Experimental source: adult testis; clone DKFZp434C192
A:Note: DKFZp434C192.1

Query Match
Best Local Similarity 86.0%; Score 51; DB 2; Length 299;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 6 PMPMPM 12
DB 37 PMPMPM 43

RESULT 4
B72492
hypothetical protein APE2577 - Aeropyrum pernix (strain K1)
C:Species: Aeropyrum pernix
C>Date: 20-Aug-1999 #sequence_revision 20-Aug-1999 #text_change 20-Aug-1999
C:Accession: B72492
R:Kawarabayashi, Y.; Hino, Y.; Horikawa, H.; Yamazaki, S.; Haikawa, Y.; Jin-no, K.; Takahara, H.; Takamiya, M.; Masuda, S.; Funahashi, T.; Tanaka, T.; Kudoh, Y.; Yamazaki, J.; K
DNA Res. 6, 83-101, 1999
A:Title: Complete genome sequence of an aerobic hyper-thermophilic Crenarchaeon, Aeropyr
A:Reference number: A72450; MUID:99310339

A:Accession: B72492
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-298 <KAW>
A:Cross-references: DDBJ:AP000064; NID:95105945; PIDN:BAA81594.1; PID:d1045380; PID:9
A:Experimental source: strain K1
C:Genetics:
A:Gene: APE2577

Query Match
Best Local Similarity 53.8%; Score 49; DB 2; Length 298;
Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

OY 2 LKMPMPM 11
DB 102 LKMPMPM 111

RESULT 5
JT0751
ferredoxin--NADP+ reductase (EC 1.18.1.2), long form precursor - bovine
M:Alternate names: adrenodoxin reductase
C:Species: Bos primigenius taurus (cattle)
C>Date: 14-Jul-1994 #sequence_revision 18-Oct-1996 #text_change 16-Jun-2000
C:Accession: JT0751; JT0079; JS0390; S03558; PS0003; A29604; S52100
R:Taketa, Y.; Sagara, Y.; Kono, A.; Sekimizu, K.; Horuchi, T.
Biol. Pharm. Bull. 16, 1200-1206, 1993
A:Title: Gene structure of bovine adrenodoxin reductase.
A:Reference number: JT0751; MUID:94177140
A:Accession: JT0751
A:Molecule type: DNA
A:Residues: 1-498 <TAK>
A:Cross-references: GB:D83475; NID:g1199916; PIDN:BAA11921.1; PID:g4521308
A:Experimental source: adrenal cortex
A:Note: the authors translated the codon GTC for residue 205 as Gly
R:Sagara, Y.; Taketa, Y.; Miyata, T.; Hara, T.; Horuchi, T.
J. Biochem. 102, 1333-1336, 1987
A:Title: Cloning and sequence analysis of adrenodoxin reductase cDNA from bovine adre
A:Reference number: JT0079; MUID:88198050
A:Accession: JT0079

A:Molecule type: mRNA
A:Residues: 1-204,211-498 <SAG>
A:Cross-references: GB:D00211; NID:g217433; PIDN:BAA00150.1; PID:g217434
A:Note: the deduced sequence is partially confirmed by amino acid sequencing of 15 is
R:Sagara, Y.
submitted to DDBJ, September 1989
A:Reference number: JS0390
A:Contents: revision, insertion of residues 205-210
A:Accession: JS0390
A:Molecule type: mRNA
A:Residues: 56-498 <SAG>
R:Hanukoglu, I.; Gutfinger, T.
Eur. J. Biochem. 180, 479-484, 1989
A:Title: cDNA sequence of adrenodoxin reductase. Identification of NADP-binding sites
A:Reference number: S03558; MUID:89170752
A:Accession: S03558
A:Molecule type: mRNA
A:Residues: 155-204,211-498 <HAN>
A:Cross-references: EMBL:X13736; NID:965; PIDN:CAA32002.1; PID:g833776
A:Note: 40S-Ser was also found
R:Hamamoto, I.; Kurokouchi, K.; Tanaka, S.; Ichikawa, Y.
Biochim. Biophys. Acta 953, 207-213, 1988
A:Title: Adrenoferradoxin-binding peptide of NADPH-adrenoferradoxin reductase.
A:Reference number: PS0003; MUID:88184054
A:Accession: PS0003

A:Molecule type: protein
A:Residues: 33-41,'S',43-62,260-283,'TM',496-498 <HAM>
A:Note: a cyanogen bromide peptide binds to adrenoferradoxin
R:Nonaka, Y.; Murakami, H.; Yabasaki, Y.; Kuramitsu, S.; Kagamiyama, H.; Yamano, T.;
Biochem. Biophys. Res. Commun. 145, 1239-1247, 1987
A:Title: Molecular cloning and sequence analysis of full-length cDNA for mRNA of adre
A:Reference number: A29604; MUID:87270696

A:Accession: A29604
 A:Molecule type: mRNA
 A:Residues: 1-76; 'R', 78-80, 'VMALTPRSMIL', 95-123, 'RVYPLT', 129-204, 211-273, 'R', 275-322,
 A:Cross-references: GB:M17029; NID:9162628; PIDN:AAA30362.1; PID:9162629
 A:Experimental source: adrenal cortex
 R:Wardburton, R.J.; Seybert, D.W.
 Biochim. Biophys. Acta 1246, 39-46, 1995
 A:Title: Structural and functional characterization of bovine adrenodoxin reductase by 1
 A:Reference number: S52100; MUID:95110846
 A:Accession: S52100
 A:Status: preliminary
 A:Molecule type: protein
 A:Residues: 'X', 34-41, 'X', 43-48, 'X', 50-51, 304-306, 'X', 308-309, 'X', 311-326 <MAR>
 C:Comment: Ferredoxin-NADP+ reductase is localized in the matrix of adrenal cortex mito
 ferredoxin-NADP+ reductase, adrenodoxin and two forms of cytochrome P-450.
 C:Genetics:
 A:Insertions: 27/1; 59/3; 91/3; 132/3; 170/3; 204/3; 246/3; 275/1; 341/3; 399/1; 456/1
 C:Function:
 A:Description: catalyzes the reversible reduction of NADP+ by reduced ferredoxin or redu
 C:Superfamily: human ferredoxin-NADP+ reductase
 C:Keywords: alternative splicing; flavoprotein; mitochondrion; monomer; NADP; oxidoreduc
 F:1-37/Domain: transit peptide (mitochondrion) #status predicted <SIG>
 F:33-498/Product: ferredoxin-NADP+ reductase, long form #status predicted <MAR>
 F:33-204, 211-498/Product: ferredoxin-NADP+ reductase, short form #status experimental <
 F:40-70/Region: beta-alpha-beta FAD nucleotide-binding fold
 F:180-190/Region: NADP binding #status predicted
 F:281/Binding site: substrate (lys) #status experimental

Query Match 53.8%; Score 49; DB 1; Length 498;
 Best Local Similarity 83.3%; Pred. No. 14;
 Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 5 WPMWPM 10
 I : : : : :
 Db 6 WPMWPM 11

RESULT 6
 S33068
 myosin heavy chain - fluke (Schistosoma mansoni) (fragment)
 N:Alternate names: surface antigen, 200K
 C:Species: Schistosoma mansoni
 C:Date: 22-Nov-1993 #sequence_revision 06-Sep-1996 #text_change 13-Feb-1998
 C:Accession: S33068
 R:Solomon, L.M.A.; Masterson, C.P.; Tom, T.D.; McNally, M.T.; Lowell, G.H.; Strand, M.
 J. Immunol. 149, 3612-3620, 1992
 A:Title: Induction of protective immunity in mice using a 62-kDa recombinant fragment of
 A:Reference number: A46514; MUID:93056536
 A:Accession: S33068
 A:Molecule type: mRNA
 A:Residues: 1-527 <SOIT>
 A:Cross-references: EMBL:X65591
 A:Note: the authors translated the codon CAA for residue 346 as Lys
 C:Superfamily: myosin heavy chain; myosin motor domain homology
 C:Keywords: ATP; surface antigen

Query Match 53.8%; Score 49; DB 2; Length 527;
 Best Local Similarity 62.5%; Pred. No. 15;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 1 ILKKMPWM 8
 I : : : : :
 Db 106 VLKMPWM 113

RESULT 7
 B70741
 probable moey protein - Mycobacterium tuberculosis (strain H37RV)
 C:Species: Mycobacterium tuberculosis
 C:Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 22-Oct-1999
 C:Accession: B70741

R:Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon
 ; Connor, R.; Davies, R.; Devlin, K.; Felwell, T.; Gentles, S.; Hamlin, N.; Holroyd,
 Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.
 Nature 393, 537-544, 1998
 A:Authors: Squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.
 A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete geno
 A:Reference number: A70500; MUID:98295987
 A:Accession: B70741
 A:Status: preliminary; nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-715 <COH>
 A:Cross-references: GB:Z75555; GB:AL123456; NID:93261608; PIDN:CAA99988.1; PID:e25035
 A:Experimental source: strain H37RV
 C:Genetics:
 A:Gene: moey

Query Match 53.8%; Score 49; DB 2; Length 715;
 Best Local Similarity 60.0%; Pred. No. 20;
 Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

OY 3 KKMPMPMR 12
 I : : : : :
 Db 64 KRWVYPMRR 73

RESULT 8
 A59287
 myosin heavy chain - fluke (Schistosoma mansoni) (strain Brazilian LE)
 C:Species: Schistosoma mansoni
 C:Date: 09-Jun-2000 #sequence_revision 09-Jun-2000 #text_change 08-Sep-2000
 C:Accession: A59287
 R:Weston, D.S.; Schmitz, J.; Kemp, M.; Kunz, W.
 Mol. Biochem. Parasitol. 58, 161-164, 1993
 A:Title: Cloning and sequence characterization of a complete myosin heavy chain cDNA
 A:Reference number: A59287; MUID:93211444
 A:Accession: A59287
 A:Status: preliminary; not compared with conceptual translation
 A:Molecule type: mRNA
 A:Residues: 1-1940 <WES>
 A:Cross-references: GB:L01634; PIDN:AAA29905.1
 A:Experimental source: strain Brazilian LE
 C:Genetics:
 A:Gene: MYH
 C:Superfamily: myosin heavy chain; myosin motor domain homology
 F:82-752/Domain: myosin motor domain homology <MMO>

Query Match 53.8%; Score 49; DB 2; Length 1940;
 Best Local Similarity 62.5%; Pred. No. 53;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 1 ILKKMPWM 8
 I : : : : :
 Db 809 VLKMPWM 816

RESULT 9
 T29295
 hypothetical protein C50F7.8 - Caenorhabditis elegans
 C:Species: Caenorhabditis elegans
 C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999
 C:Accession: T29295
 R:Johnson, D.; Stellyes, L.
 submitted to the EMBL Data Library, November 1995
 A:Description: The sequence of C. elegans cosmid C50F7.
 A:Reference number: Z20601
 A:Accession: T29295
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-111 <COH>
 A:Cross-references: EMBL:U41557; PIDN:AAA83303.1; CESP:C50F7.8
 C:Genetics:

C:Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 20-Sep-1999
 C:Accession: S37664
 R:Kusters, J.G.; Jager, E.J.; Niesters, H.G.M.; van der Zeijst, B.A.M.
 Vaccine 8, 605-608, 1990
 A:Title: Sequence evidence for RNA recombination in field isolates of avian coronavirus
 A:Reference number: S37663; MUID:91205880
 A:Accession: S37664
 A:Molecule type: genomic RNA
 A:Residues: 1-621 <KUS>
 A:Cross-references: EMBL:X58001; NID:g58986; PIDN:CAA41065.1; PID:g58987
 C:Superfamily: coronavirus E2 glycoprotein
 C:Keywords: glycoprotein; peplomer protein; spike protein
 F:1-5/Product: E2 glycoprotein subunit S1 (fragment) #status predicted <GS1>
 F:6-621/Product: E2 glycoprotein subunit S2 #status predicted <GS2>
 F:10,47,59,137,144,415,447,482,506,519,542/Binding site: carbohydrate (Asn) (covalent) #

Query Match 51.1%; Score 46.5; DB 2; Length 621;
 Best Local Similarity 61.5%; Pred. No. 38;
 Matches 8; Conservative 1; Mismatches 1; Indels 3; Gaps 1;

OY 1 ILK--KMPMPW 10
 ||| ||||: |
 Db 553 ILKTYIKMPWYVW 565

RESULT 15

S37663
 Peplomeric polyprotein precursor - avian infectious bronchitis virus (strain D207) (frag
 N:Contains: E2 glycoprotein subunit S2
 C:Species: avian infectious bronchitis virus, IBV
 A:Variety: strain D207
 C:Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 20-Sep-1999
 C:Accession: S37663
 R:Kusters, J.G.; Jager, E.J.; Niesters, H.G.M.; van der Zeijst, B.A.M.
 Vaccine 8, 605-608, 1990
 A:Title: Sequence evidence for RNA recombination in field isolates of avian coronavirus
 A:Reference number: S37663; MUID:91205880
 A:Accession: S37663
 A:Molecule type: genomic RNA
 A:Residues: 1-630 <KUS>
 A:Cross-references: EMBL:X58003; NID:g58988; PIDN:CAA41067.1; PID:g58989
 C:Superfamily: coronavirus E2 glycoprotein
 C:Keywords: glycoprotein; peplomer protein; spike protein
 F:1-5/Product: E2 glycoprotein subunit S1 (fragment) #status predicted <GS1>
 F:6-621/Product: E2 glycoprotein subunit S2 #status predicted <GS2>
 F:10,47,59,137,144,415,447,482,506,519,542/Binding site: carbohydrate (Asn) (covalent) #

Query Match 51.1%; Score 46.5; DB 2; Length 630;
 Best Local Similarity 61.5%; Pred. No. 39;
 Matches 8; Conservative 1; Mismatches 1; Indels 3; Gaps 1;

OY 1 ILK--KMPMPW 10
 ||| ||||: |
 Db 553 ILKTYIKMPWYVW 565

Search completed: January 4, 2002, 08:41:32
 Job time: 170 sec

THIS PAGE BLANK (USPTO)

AUTHORS Sasaki, T. and Yamamoto, K.
 TITLE Rice cDNA from green shoot (2001)
 JOURNAL Unpublished (2001)
 COMMENT Contact: Takuji Sasaki
 National Institute of Agrobiological Resources
 Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki
 305-8602, Japan
 Tel: 81-298-38-7441
 Fax: 81-298-38-7468
 Email: tsasaki@agr.affrc.go.jp, URL: http://rgrp.dna.affrc.go.jp/
 PROJECT = 'RGP',
 S16019_97A.

FEATURES

source Location/Qualifiers
 1. 448

/organism="Oryza sativa"
 /strain="Nipponbare"
 /db_xref="taxon:4530"
 /clone="S16019"
 /clone_lib="Rice green shoot"
 /note="Green shoot (8 days old)"

BASE COUNT 85 a 146 c 160 g 57 t
 ORIGIN

alignment_scores:

Quality: 69.00 Length: 10
 Ratio: 7.667 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:

US-09-444-281-36 x AU198162/rev ..

Align seg 1/1 to reverse of: AU198162 from: 1 to: 448

3 ArgTrpProTrrPrProTrrPArgArgLys 12
 |||||
 311 CCCTGCGCTTGCTGCGCCGCGCGCGG 282

seq_name: gb_est1:AU082117

seq_documentation_block:

LOCUS AU082117 578 bp mRNA EST 04-FEB-2000
 DEFINITION AU082117 Rice panicle at ripening stage Oryza sativa cDNA clone
 E11611, mRNA sequence.

ACCESSION AU082117

VERSION AU082117.1 GI:6727452

KEYWORDS EST.

SOURCE Oryza sativa.

ORGANISM Oryza sativa

REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 Ehrhartoideae; Oryzoideae; Oryza.

AUTHORS Sasaki, T. and Yamamoto, K.

TITLE Rice cDNA from panicle at ripening stage (2000)

JOURNAL Unpublished (2000)

COMMENT Contact: Takuji Sasaki
 National Institute of Agrobiological Resources
 Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki
 305-8602, Japan
 Tel: 81-298-38-7441
 Fax: 81-298-38-7468
 Email: tsasaki@agr.affrc.go.jp, URL: http://rgrp.dna.affrc.go.jp/
 PROJECT = 'RGP',

FEATURES

source

1. 578 Location/Qualifiers

/organism="Oryza sativa"
 /strain="Nipponbare"
 /db_xref="taxon:4530"
 /clone="E11611"
 /clone_lib="Rice panicle at ripening stage"
 /dev_stage="ripening stage"
 /note="Organ: panicle; Rice cDNA from panicle at ripening

BASE COUNT 129 a 180 c 179 g 90 t
 ORIGIN

alignment_scores:

Quality: 69.00 Length: 10
 Ratio: 7.667 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:

US-09-444-281-36 x AU082117/rev ..

Align seg 1/1 to reverse of: AU082117 from: 1 to: 578

3 ArgTrpProTrrPrProTrrPArgArgLys 12
 |||||
 345 CCCTGCGCTTGCTGCGCCGCGCGCGG 316

seq_name: gb_est1:AU089922

seq_documentation_block:

LOCUS AU089922 446 bp mRNA EST 19-APR-2000
 DEFINITION AU089922 Hordeum vulgare subsp. vulgare upper three leaves at
 heading stage Hordeum vulgare subsp. vulgare cDNA clone
 haruna_1lib1_121, mRNA sequence.

ACCESSION AU089922

VERSION AU089922.1 GI:7613350

KEYWORDS EST.

SOURCE Hordeum vulgare subsp. vulgare.

ORGANISM Hordeum vulgare subsp. vulgare

REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooidae
 ; Triticeae; Hordeum.

AUTHORS Sato, K., Takahashi, H. and Takeda, K.

TITLE Hordeum vulgare subsp. vulgare cDNA clone

JOURNAL Unpublished (2000)

COMMENT Contact: Kazuhiko Sato
 Research Institute for Bioreources
 Okayama University, Barley Germplasm Center
 Chuo 2-20-1, Kurashiki, Okayama 710-0046, Japan
 Email: kazsato@rib.okayama-u.ac.jp
 URL: http://www.rib.okayama-u.ac.jp/barley/.

Location/Qualifiers

1. 446

/organism="Hordeum vulgare subsp. vulgare"

/cultivar="Haruna N130"

/db_xref="taxon:112509"

/clone="haruna_1lib1_121"

/clone_lib="Hordeum vulgare subsp. vulgare upper three
 leaves at heading stage"

/tissue_type="upper three leaves at heading stage"

BASE COUNT 89 a 130 c 149 g 76 t 2 others

ORIGIN

alignment_scores:

Quality: 68.00 Length: 8
 Ratio: 8.500 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-36 x AU089922/rev ..

Align seg 1/1 to reverse of: AU089922 from: 1 to: 446

4 TrpProTrrPrProTrrPArgArg 11
 |||||
 188 TGCGGTGCTGCGCGCGCGG 165

seq_name: gb_est1:AU089934

seq_documentation_block: 475 bp mRNA EST 19-Apr-2000
 LOCUS AU089934 Hordeum vulgare subsp. vulgare upper three leaves at
 DEFINITION AU089934 Hordeum vulgare subsp. vulgare upper three leaves at
 heading stage Hordeum vulgare subsp. vulgare cDNA clone
 haruna_11b1_134, mRNA sequence.
 accession AU089934
 version AU089934.1 GI:7613362
 source EST
 ORGANISM Hordeum vulgare subsp. vulgare.
 Hordeum vulgare subsp. vulgare.
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooidae
 ; Triticeae; Hordeum.
 1 (bases 1 to 475)
 Sato, K., Takahashi, H. and Takeda, K.
 Hordeum vulgare subsp. vulgare cDNA clone
 Unpublished (2000)
 Contact: Kazuhiro Sato
 Research Institute for Bioresearches
 Okayama University, Barley Germplasm Center
 Chuo 2-20-1, Kurashiki, Okayama 710-0046, Japan
 Email: kazsato@rib.okayama-u.ac.jp,
 URL: http://www.rib.okayama-u.ac.jp/barley/
 Location/Qualifiers
 1..475
 /organism="Hordeum vulgare subsp. vulgare"
 /cultivar="Haruna Nijo"
 /db_xref="taxon:112509"
 /clone="haruna_11b1_134"
 /clone_1lb="Hordeum vulgare subsp. vulgare upper three
 leaves at heading stage"
 /tissue_type="Upper three leaves at heading stage"
 BASE COUNT 100 a 137 c 155 g 83 t
 ORIGIN

alignment_scores:
 Quality: 68.00 Length: 8
 Ratio: 8.500 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
 US-09-444-281-36 x AU089934/rev ..

Align seg 1/1 to reverse of: AU089934 from: 1 to: 475

4 TrpProTrrTrrProTrrPargarg 11
 |||
 194 TGGCGTGGTGGCGCGCGCGCGA 171

seq_name: gb_est1:BE024584

seq_documentation_block:
 LOCUS BE024584 330 bp mRNA EST 06-JUN-2000
 DEFINITION 894003H02.y1 C. reinhardtii CC-1690, normalized, lambda zap II
 Chlamydomonas reinhardtii cDNA, mRNA sequence.
 accession BE024584
 version BE024584.1 GI:8287025
 source EST
 ORGANISM Chlamydomonas reinhardtii.
 Chlamydomonas reinhardtii.
 Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
 Chlamydomonadaceae; Chlamydomonas.
 1 (bases 1 to 330)
 Grossman, A., Davies, J., Federspiel, N., Harris, E., Lefebvre, P.,
 Mcdermott, J. P., Sillflow, C., Stern, D. and Surzycki, R.
 Analyses of the Chlamydomonas reinhardtii Genome: A Model,
 Unicellular System for Analyzing Gene Function and Regulation in
 Vascular Plants: project phase 2
 Unpublished (2000)
 Contact: Elizabeth H. Harris
 DCMB Box 91000
 Duke University

JOURNAL
 COMMENT

Durham, NC 27708-1000, USA
 Tel: 919 613 8164
 Fax: 919 613 8177
 Email: chlamy@duke.edu.
 Location/Qualifiers
 1..330
 /organism="Chlamydomonas reinhardtii"
 /strain="CC-1690 wild type mt+ 21gr"
 /db_xref="taxon:3055"
 /clone_1lb="C. reinhardtii CC-1690, normalized, lambda zap
 II"
 /note="Vector: pBluescript II SK-. Site 1: EcoRI; Site 2:
 XhoI; This library, constructed by John Davies and Jeffrey
 Mcdermott, combines cDNAs from CC-1690 cells grown to
 mid-log phase in TAP (acetate-containing) medium in the
 light, TAP medium in the dark, HS (minimal) medium in
 ambient levels of CO2 and HS medium bubbled with 5% CO2.
 PolyA mRNA was purified from each sample, pooled and cDNA
 synthesized. The cDNA was directionally cloned into lambda
 ZAP II (Stratagene) in the EcoRI (5') and XhoI (3') sites.
 pBluescript II SK- plasmids were excised from the lambda
 ZAP clones by superinfection with Exsist (Stratagene)
 phage. The library was normalized using method 4 described
 in Bonaldo et al (1996) Genome Research 6: 791-806."

BASE COUNT 40 a 103 c 142 g 45 t
 ORIGIN

alignment_scores:
 Quality: 67.00 Length: 9
 Ratio: 8.375 Gaps: 0
 Percent Similarity: 88.889 Percent Identity: 88.889

alignment_block:
 US-09-444-281-36 x BE024584 ..

Align seg 1/1 to: BE024584 from: 1 to: 330

3 ArgTrpProTrrTrrProTrrPargarg 11
 |||
 72 CGGTGGCGTGGTGGCGCGCGG 98

seq_name: gb_est1:AUI98258

seq_documentation_block:
 LOCUS AUI98258 352 bp mRNA EST 12-JUL-2001
 DEFINITION AUI98258 Rice green shoot Oryza sativa cDNA clone S16389, mRNA
 sequence.
 accession AUI98258
 version AUI98258.1 GI:14714335
 source EST
 ORGANISM Oryza sativa.
 Oryza sativa.
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 Ehrhartoidae; Oryzaceae; Oryza.
 1 (bases 1 to 352)
 Sasaki, T. and Yamamoto, K.
 Rice cDNA from green shoot (2001)
 Unpublished (2001)
 Contact: Takuji Sasaki
 National Institute of Agrobiological Resources
 Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki
 305-8602, Japan
 Tel: 81-298-38-7441
 Fax: 81-298-38-7468
 Email: tsasaki@br.affrc.go.jp, URL: http://rgp.dna.affrc.go.jp/
 PROJECT = 'RGP'
 S16389_967.

REFERENCE
 AUTHORS
 TITLE
 JOURNAL
 COMMENT

FEATURES
 source
 1..352
 /organism="Oryza sativa"
 /strain="Nipponbare"

/db_xref="taxon:4530"
/clone="S16389"
/clone_lib="Rice green shoot"
/note="Green shoot (8 days old)"
BASE COUNT 95 a 95 c 104 g 51 t 7 others
ORIGIN

alignment_scores:
Quality: 67.00 Length: 9
Ratio: 8.375 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 88.889

alignment_block:
US-09-444-281-36 x AV61634/rev ..

Align seg 1/1 to reverse of: AU198258 from: 1 to: 352

3 ArgTrrProTrrPrrProTrrPrrArg 11
|||||
167 CGCTGGCGCTGTGGTGGCGGCGG 141

seq_name: gb_est1:AV61634

seq_documentation_block:
LOCUS AV61634 371 bp mRNA EST 15-DEC-2000
DEFINITION AV61634 Chlamydomonas reinhardtii 5% CO2 Chlamydomonas reinhardtii
ACCESSION AV61634
VERSION AV61634.1 GI:10784962
KEYWORDS EST.
SOURCE Chlamydomonas reinhardtii.
ORGANISM Chlamydomonas reinhardtii
Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
Chlamydomonadales; Chlamydomonas.

REFERENCE 1 (bases 1 to 371)
Asanizu,E., Miura,K., Kuchio,K., Inoue,Y., Fukuzawa,H., Ohyama,K.,
Nakamura,Y. and Tabata,S.
Generation of expressed sequence tags from low-CO2 and high-CO2
adapted cells of Chlamydomonas reinhardtii
DNA Res. 7 (5), 805-307 (2000)
20539644

JOURNAL MEDLINE
COMMENT Contact: Erika Asanizu
The First Laboratory for Plant Gene Research
Kazusa DNA Research Institute
Yana 1532-3, Kisarazu, Chiba 292-0812, Japan
Email: asanizuekazusa.or.jp. URL: http://www.kazusa.or.jp/en/plant/.

FEATURES
source Location/Qualifiers
1..371
/organism="Chlamydomonas reinhardtii"
/strain="C9"
/db_xref="taxon:3055"
/clone_lib="HCL037H03.F"

/note="Vector: Bluescript II SK-; Site_1: EcoRI; Site_2:
XhoI; The cDNA library was constructed from cells cultured
in a medium with bubbling air containing 5% carbon
dioxide"

BASE COUNT 73 a 106 c 141 g 51 t
ORIGIN

alignment_scores:
Quality: 67.00 Length: 9
Ratio: 8.375 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 88.889

alignment_block:

US-09-444-281-36 x AV61634 ..

Align seg 1/1 to: AV61634 from: 1 to: 371

3 ArgTrrProTrrPrrProTrrPrrArg 11

|||||
244 CGGTGGCGCTGTGGCGGCGGCGG 270

seq_name: gb_est1:BE129188

seq_documentation_block:
LOCUS BE129188 386 bp mRNA EST 21-JUN-2000
DEFINITION BE129188 y1 C. reinhardtii CC-1690, normalized, Lambda zap II
Chlamydomonas reinhardtii cDNA, mRNA sequence.

ACCESSION BE129188
VERSION BE129188.1 GI:8576551

KEYWORDS EST.
SOURCE Chlamydomonas reinhardtii.
ORGANISM Chlamydomonas reinhardtii
Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
Chlamydomonadales; Chlamydomonas.

REFERENCE 1 (bases 1 to 386)
Grossman,A., Davies,J., Federspiel,N., Harris,E., Lefebvre,P.,
McDermott,J.P., Sillow,C., Stern,D. and Surzycki,R.
Analyses of the Chlamydomonas reinhardtii Genome: A Model,
Unicellular System for Analyzing Gene Function and Regulation in
Vascular Plants: project phase 2

JOURNAL

COMMENT Unpublished (2000)
Contact: Elizabeth H. Harris
DCMB Box 91000
Duke University
Durham, NC 27708-1000, USA
Tel: 919 613 8164
Fax: 919 613 8177
Email: chlamy@duke.edu.

FEATURES
source Location/Qualifiers
1..386
/organism="Chlamydomonas reinhardtii"
/strain="CC-1690 wild type mt+ 21gr"
/db_xref="taxon:3055"
/clone_lib="C. reinhardtii CC-1690, normalized, Lambda zap
II"

/note="Vector: pBluescript II SK-; Site_1: EcoRI; Site_2:
XhoI; This library, constructed by John Davies and Jeffrey
McDermott, combines cDNAs from CC-1690 cells grown to
mid-log phase in TAP (acetate-containing) medium in the
light, TAP medium in the dark, HS (minimal) medium in
ambient levels of CO2 and HS medium bubbled with 5% CO2.
Polya mRNA was purified from each sample, pooled and cDNA
synthesized. The cDNA was directionally cloned into lambda
zap II (Stratagene) in the EcoRI (5') and XhoI (3') sites.
pBluescript II SK- plasmids were excised from the lambda
zap clones by superinfection with EXAssist (Stratagene)
phage. The library was normalized using method 4 described
in Bonaldo et al (1996) Genome Research 6: 791-806."

BASE COUNT 79 a 109 c 142 g 56 t
ORIGIN

alignment_scores:
Quality: 67.00 Length: 9
Ratio: 8.375 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 88.889

alignment_block:
US-09-444-281-36 x BE129188 ..

Align seg 1/1 to: BE129188 from: 1 to: 386

3 ArgTrrProTrrPrrProTrrPrrArg 11
|||||
249 CGGTGGCGCTGTGGCGGCGGCGG 275

seq_name: gb_est2:BF584442

seq_documentation_block:
LOCUS BF584442 701 bp mRNA EST 12-DEC-2000
DEFINITION 602098336F1 NCL_CGAP_Co24 Mus musculus cDNA clone IMAGE:4218273 5',

ACCESSION mRNA sequence.
BF584442
VERSION BF584442.1 GI:11658160
KEYWORDS EST.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 701)
TITLE NIH-MGC http://mgc.nci.nih.gov/
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabs@mail.nih.gov
Tissue Procurement: Jeffrey E. Green, M.D.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at: <http://image.llnl.gov>
Plate: LAM9798 row: g column: 10
High quality sequence stop: 701.

FEATURES
source
1..701
/organism="Mus musculus"
/strain="FVB/N"
/db_xref="taxon:10090"
/clone="IMAGE:4218273"
/lab_host="DH10B (T1 phage-resistant)"
/note="Organ: colon; Vector: PCMV-SpOrf; Site: 1: NotI; Site: 2: SalI; Cloned unidirectionally. Primer: Oligo dT. Average insert size 1.6 kb. Constructed by Life Technologies. Note: this is a NCI_CGAP library."

BASE COUNT 149 a 198 c 186 g 168 t
ORIGIN

alignment_scores:
Quality: 67.00 Length: 10
Ratio: 7.444 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:
US-09-444-281-36 x BF584442
Align seg 1/1 to: BF584442 from: 1 to: 701
11leuAgtTrrpTrpTrpTrpArg 10
:::||||| ||||||| ||||||| |||||||
188 CTCTTATTGTGCGCTTGTGCGCATGAGCA 217

seq_name: gb_est2:BG964576

seq_documentation_block:
LOCUS BG964576 834 bp mRNA EST 12-JUN-2001
DEFINITION 6028322551 NCI_CGAP_CO24 Mus musculus cDNA clone IMAGE:4966828 5',
mRNA sequence.
ACCESSION BG964576
VERSION BG964576.1 GI:14352213
KEYWORDS EST.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 834)
TITLE NIH-MGC <http://mgc.nci.nih.gov/>.
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabs@mail.nih.gov
Tissue Procurement: Jeffrey E. Green, M.D.

CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at: <http://image.llnl.gov>
Plate: LAM10996 row: n column: 13
High quality sequence stop: 811.

FEATURES
source
1..834
/organism="Mus musculus"
/strain="FVB/N"
/db_xref="taxon:10090"
/clone="IMAGE:496828"
/lab_host="DH10B (T1 phage-resistant)"
/note="Organ: colon; Vector: PCMV-SpOrf; Site: 1: NotI; Site: 2: SalI; Cloned unidirectionally. Primer: Oligo dT. Average insert size 1.6 kb. Constructed by Life Technologies. Note: this is a NCI_CGAP library."

BASE COUNT 185 a 202 c 226 g 221 t
ORIGIN

alignment_scores:
Quality: 67.00 Length: 10
Ratio: 7.444 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:
US-09-444-281-36 x BG964576
Align seg 1/1 to: BG964576 from: 1 to: 834
11leuAgtTrrpTrpTrpTrpArg 10
:::||||| ||||||| ||||||| |||||||
195 CTGTTATTGTGCGCTTGTGCGCATGAGCA 224

seq_name: gb_gss:A0943724

seq_documentation_block:
LOCUS A0943724 514 bp DNA GSS 27-JAN-2000
DEFINITION Sheared DNA-36H6.TF Sheared DNA Trypanosoma brucei genomic clone
ACCESSION A0943724
VERSION A0943724.1 GI:6766989
KEYWORDS GSS.
SOURCE Trypanosoma brucei.
ORGANISM Trypanosoma brucei
Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;
Trypanosoma.
REFERENCE 1 (bases 1 to 514)
TITLE El-Sayed, N., Zhao, S., Zhao, H., Gill, S., Suh, E., Malek, J., Fujii, C., Gerrard, C., Leech, V., de Jong, P., Ullu, E., Melville, S., Donelson, J., Fraser, C. and Adams, M.
AUTHORS Determination of clone end sequences from Trypanosoma brucei cDNA
JOURNAL 10.1 sheared DNA library
COMMENT Other_GSS: Sheared DNA-36H6.TF
Unpublished (1999)
Contact: Najib M. El-Sayed
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: nelsayed@igf.org
Clones are derived from the Trypanosoma brucei cDNA 10.1 sheared DNA library constructed at TIGR. Clones will be available for distribution through ATCC. Sheared DNA end sequences search page: <http://www.tigr.org/tdb/mdb/cdb/>.
Seq primer: M13-forward
Class: shotgun.
Location/Qualifiers

source
1. .514
/organism="Trypanosoma brucei"
/strain="TREU927/4 GUTat 10.1"
/db_xref="taxon:5691"
/clone="Sheared DNA-36H6"
/note="Vector: pUC18; Site_1: Small; Constructed at The Institute for Genomic Research (TIGR), Rockville, MD. Genomic DNA isolated from a cloned population of Trypanosoma brucei (TREU927/4 GUTat 10.1) was mechanically sheared to give a tight size distribution (approx 2 kb). The v + i method used for the library construction is described in detail in Smith, H.O. and Venter, J.C. (Making small insert libraries for whole genome shotgun sequencing projects. In Genome Sequencing: A Practical Approach, eds. M. Vaubin and B. Barrell, Oxford University Press, 1999)."

BASE COUNT 124 a 185 c 131 g 74 t
ORIGIN

alignment_scores:
Quality: 66.00 Length: 10
Ratio: 7.333 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block:
US-09-444-281-36 x AQ943724/rev ..
Align seg 1/1 to reverse of: AQ943724 from: 1 to: 514

2 LeuArgTPrProTPrProTPrArg 11
|||||
450 CTTCGATGGCCTTGCTGTGGTGGCGCGG 421

seq_name: gb_gss:AZ218599
seq_documentation_block:
LOCUS AZ218599 512 bp DNA GSS 09-JUN-2000
DEFINITION Sheared DNA-82B1.TR Sheared DNA Trypanosoma brucei genomic clone
ACCESSION AZ218599
VERSION AZ218599.1 GI:8436399
KEYWORDS GSS.
SOURCE Trypanosoma brucei.
ORGANISM Trypanosoma brucei.
Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;
Trypanosoma.
1 (bases 1 to 572)
El-Sayed, N., Zhao, S., Zhao, H., Gill, S., Suh, E., Malek, J., Fujii, C.,
Gerrard, C., Leech, V., de Jong, P., Ullu, E., Melville, S., Donelson, J.,
Fraser, C. and Adams, M.
Determination of clone end sequences from Trypanosoma brucei GUTat
10.1 sheared DNA library
Unpublished (1999)
Other GSSs: Sheared DNA-82B1.TF
Contact: Najib M. El-Sayed
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: nelsayed@tigr.org
Clones are derived from the Trypanosoma brucei GUTat 10.1 sheared
DNA library constructed at TIGR. Clones will be available for
distribution through Research Genetics, Alabama, USA. Sheared DNA
end sequences search page: <http://www.tigr.org/tdb/mbd/tbdt/>.
Seq primer: M13-Reverse
Class: shotgun.
FEATURES
location/Qualifiers
1. .572
/organism="Trypanosoma brucei"
/strain="TREU927/4 GUTat 10.1"

/db_xref="taxon:5691"
/clone="Sheared DNA-82B1"
/clone_lib="Sheared DNA"
/note="Vector: pUC18; Site_1: Small; Constructed at The Institute for Genomic Research (TIGR), Rockville, MD. Genomic DNA isolated from a cloned population of Trypanosoma brucei (TREU927/4 GUTat 10.1) was mechanically sheared to give a tight size distribution (approx 2 kb). The v + i method used for the library construction is described in detail in Smith, H.O. and Venter, J.C. (Making small insert libraries for whole genome shotgun sequencing projects. In Genome Sequencing: A Practical Approach, eds. M. Vaubin and B. Barrell, Oxford University Press, 1999)."

BASE COUNT 136 a 217 c 144 g 75 t
ORIGIN

alignment_scores:
Quality: 66.00 Length: 10
Ratio: 7.333 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block:
US-09-444-281-36 x AZ218599/rev ..
Align seg 1/1 to reverse of: AZ218599 from: 1 to: 572

2 LeuArgTPrProTPrProTPrArg 11
|||||
211 CTTCGATGGCCTTGCTGTGGTGGCGCGG 182

seq_name: gb_est2:C52835
seq_documentation_block:
LOCUS C52835 208 bp mRNA EST 11-SEP-1997
DEFINITION C52835 yuji Kohara unpublished cdNA Caenorhabditis elegans cdNA
clone yK285b1 3', mRNA sequence.
ACCESSION C52835
VERSION C52835.1 GI:2390592
KEYWORDS EST.
SOURCE Caenorhabditis elegans.
ORGANISM Caenorhabditis elegans.
Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea
; Rhabditidae; Peloderinae; Caenorhabditis.
1 (bases 1 to 208)
Kohara, Y., Morishashi, T., Tabara, H., Watanabe, H., Sugimoto, A., Sano
M., Miyata, A. and Nishigaki, A.
Expression map of the C. elegans genome
Unpublished (1996)
Contact: Yuji Kohara
Genome Biology Lab.
National Institute of Genetics
Yata 1111, Mishima, Shizuoka 411, Japan
Tel: 81-559-81-6854
Fax: 81-559-81-6855
Email: ykohara@lab.nig.ac.jp.
location/Qualifiers
1. .208
/organism="Caenorhabditis elegans"
/strain="CHI489 him-8(e1489)"
/db_xref="taxon:6239"
/clone="yK285b1"
/clone_lib="yuji Kohara unpublished cdNA"
/sex="hermaphrodite, male"
/tissue_type="whole animal"
/dev_stage="varied"

BASE COUNT 25 a 50 c 77 g 52 t 4 others
ORIGIN

alignment_scores:
Quality: 65.00 Length: 10

Ratio:	7.222	Gaps:	0
Percent Similarity:	90.000	Percent Identity:	70.000

alignment_block:
US-09-444-281-36 x C52835

Align seg 1/1 to: C52835 from: 1 to: 208

```

1  ILELeuArgTrpProTrpTrpProTrpArg 10
   ::::: |||||
47 GTGATGTGTGGCCCTGCTGGCCCTGGCGG 76

```

seq_name: gb_est2:C62320

```
seq_documentation_block:
  locs: C62320
```

LOCUS	292 bp	mrna	EST	22-SEP-1997
C62320				
DEFINITION	Yuji Kohara unpublished	cdna	Caenorhabditis elegans	cdna
C62320				

clone yk285b1 5', mRNA sequence.
ACCESSION G53320

ACCESSION	C62320
VERSION	C62320.1

KEYWORDS	EST.
SOURCE	Caenorhabditis elegans.

ORGANISM

Eukaryota; Metazoa; Chromadorea; Rhabdilitida; Rhabdilitoidea
; Rhabdilitidae; Pelodermineae; Caenorhabdilitis.
1 / pages 1 to 203

REFERENCE

AUTHORS Kohara, Y., Motohashi, T., Tabara, H., Watanabe, H., Sugimoto, A., Sano

TITLE Expression map of the C.elegans genome

JOURNAL
COMMENT

COMMENT
Contact: Yuji Kohara
Genome Biology Lab.

National Institute of Genetics
Yata 1111, Mishima, Shizuoka 411, Japan

Tel: 81-559-81-6854
Fax: 81-559-81-6855

Email: ykohara@lab.nig.ac.jp

FEATURES	Location/Qualifiers
source	1. .292

```
/organism="Caenorhabditis elegans"
/strain="CB1489 him-8(e1489)"
```

```
/db_xref="taxon:
/clone="yk285b1"
```

```
/clone_lib="Yuji Kohara unpublished cDNA"
```

```
/sex="hermaphrodite, male"  
/tissue_type="whole animal"
```

```

/dev_stage="varied"
BASE COUNT  94 a      89 c  60 g  49 t

```

ORIGIN

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65
66
67
68
69
70
71
72
73
74
75
76
77
78
79
80
81
82
83
84
85
86
87
88
89
90
91
92
93
94
95
96
97
98
99
100
101
102
103
104
105
106
107
108
109
110
111
112
113
114
115
116
117
118
119
120
121
122
123
124
125
126
127
128
129
130
131
132
133
134
135
136
137
138
139
140
141
142
143
144
145
146
147
148
149
150
151
152
153
154
155
156
157
158
159
160
161
162
163
164
165
166
167
168
169
170
171
172
173
174
175
176
177
178
179
180
181
182
183
184
185
186
187
188
189
190
191
192
193
194
195
196
197
198
199
200
201
202
203
204
205
206
207
208
209
210
211
212
213
214
215
216
217
218
219
220
221
222
223
224
225
226
227
228
229
230
231
232
233
234
235
236
237
238
239
240
241
242
243
244
245
246
247
248
249
250
251
252
253
254
255
256
257
258
259
260
261
262
263
264
265
266
267
268
269
270
271
272
273
274
275
276
277
278
279
280
281
282
283
284
285
286
287
288
289
290
291
292
293
294
295
296
297
298
299
300
301
302
303
304
305
306
307
308
309
310
311
312
313
314
315
316
317
318
319
320
321
322
323
324
325
326
327
328
329
330
331
332
333
334
335
336
337
338
339
340
341
342
343
344
345
346
347
348
349
350
351
352
353
354
355
356
357
358
359
360
361
362
363
364
365
366
367
368
369
370
371
372
373
374
375
376
377
378
379
380
381
382
383
384
385
386
387
388
389
390
391
392
393
394
395
396
397
398
399
400
401
402
403
404
405
406
407
408
409
410
411
412
413
414
415
416
417
418
419
420
421
422
423
424
425
426
427
428
429
430
431
432
433
434
435
436
437
438
439
440
441
442
443
444
445
446
447
448
449
450
451
452
453
454
455
456
457
458
459
460
461
462
463
464
465
466
467
468
469
470
471
472
473
474
475
476
477
478
479
480
481
482
483
484
485
486
487
488
489
490
491
492
493
494
495
496
497
498
499
500
501
502
503
504
505
506
507
508
509
510
511
512
513
514
515
516
517
518
519
520
521
522
523
524
525
526
527
528
529
530
531
532
533
534
535
536
537
538
539
540
541
542
543
544
545
546
547
548
549
550
551
552
553
554
555
556
557
558
559
560
561
562
563
564
565
566
567
568
569
570
571
572
573
574
575
576
577
578
579
580
581
582
583
584
585
586
587
588
589
590
591
592
593
594
595
596
597
598
599
600
601
602
603
604
605
606
607
608
609
610
611
612
613
614
615
616
617
618
619
620
621
622
623
624
625
626
627
628
629
630
631
632
633
634
635
636
637
638
639
640
641
642
643
644
645
646
647
648
649
650
651
652
653
654
655
656
657
658
659
660
661
662
663
664
665
666
667
668
669
670
671
672
673
674
675
676
677
678
679
680
681
682
683
684
685
686
687
688
689
690
691
692
693
694
695
696
697
698
699
700
701
702
703
704
705
706
707
708
709
710
711
712
713
714
715
716
717
718
719
720
721
722
723
724
725
726
727
728
729
730
731
732
733
734
735
736
737
738
739
740
741
742
743
744
745
746
747
748
749
750
751
752
753
754
755
756
757
758
759
760
761
762
763
764
765
766
767
768
769
770
771
772
773
774
775
776
777
778
779
780
781
782
783
784
785
786
787
788
789
790
791
792
793
794
795
796
797
798
799
800
801
802
803
804
805
806
807
808
809
810
811
812
813
814
815
816
817
818
819
820
821
822
823
824
825
826
827
828
829
830
831
832
833
834
835
836
837
838
839
840
84

```
alignment_scores: 65.00
quality: 1000
length: 1000
```

Ratio:	7.222	Gaps:	0
Percent Similarity:	90.000	Percent Identity:	70.000

```
alignment_block:
```

US-09-444-281-36 x C62320/rev ..
Align seg 1/1 to reverse of: C62320 from: 1 to: 292

```

1 IleLeuArgTrpProTrpTrpProTrpArg 10
  ::::: |||||
163 GTGATGTGTGGCCCTGTGTGGCCCTGGCGG 134

```

THIS PAGE BLANK (USPTO)

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:47:24 ; Search time 50.17 Seconds
(Without alignments)
34.986 Million cell updates/sec

Title: US-09-444-281-36
Perfect score: 86
Sequence: 1 ILRPMWPMWRK 12

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 473505 seqs, 146272329 residues

Total number of hits satisfying chosen parameters: 473505

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :
1: SPREMBL_17:*
2: sp_archaea:*
3: sp_bacteria:*
4: sp_fungi:*
5: sp_human:*
6: sp_invertebrate:*
7: sp_mhc:*
8: sp_oranelle:*
9: sp_phage:*
10: sp_plant:*
11: sp_rodent:*
12: sp_virus:*
13: sp_vertebrate:*
14: sp_unclassified:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	57	66.3	723	12 Q9DUC4	Q9DUC4 tt virus. o
2	53	61.6	746	12 Q9JH31	Q9JH31 tt virus. o
3	53	61.6	1173	12 Q990M4	Q990M4 human coron
4	53	61.6	1173	12 Q990M3	Q990M3 human coron
5	53	61.6	1173	12 Q990M2	Q990M2 human coron
6	53	61.6	1173	12 Q990M1	Q990M1 human coron
7	53	61.6	1383	12 Q84712	Q84712 porcine epi
8	52	60.5	1245	3 Q9YV5	Q9YV5 trichoderma
9	51	59.3	299	4 Q9YV1	Q9YV1 homo sapien
10	51	59.3	504	2 P96143	P96143 thermococin
11	50	58.1	111	5 Q18753	Q18753 caenorhabdi
12	50	58.1	141	1 Q9CZAI	Q9CZAI mus musculu
13	50	58.1	327	10 Q9AUN3	Q9AUN3 oryza sativ
14	50	58.1	735	12 Q9DUC9	Q9DUC9 tt virus. o
15	49	57.0	49	12 Q9D80	Q9D80 tt virus. o
16	49	57.0	226	4 Q9BS58	Q9BS58 homo sapien
17	49	57.0	467	5 Q19573	Q19573 caenorhabdi
18	49	57.0	748	12 Q9D81	Q9D81 tt virus. o
19	48.5	56.4	114	2 Q9X8C2	Q9X8C2 streptomyce

20	48	55.8	540	2	007504	007504 bacillus me
21	47	54.7	165	10	Q9SNN3	Q9SNN3 oryza sativ
22	47	54.7	276	2	Q9HXC9	Q9HXC9 pseudomonas
23	47	54.7	1411	10	Q9LYG0	Q9LYG0 arabidopsis
24	46	53.5	154	2	Q9R6J3	Q9R6J3 agrobacteri
25	46	53.5	728	3	Q9P3G0	Q9P3G0 neurospora
26	45	52.3	159	2	Q9KZT3	Q9KZT3 streptomyce
27	45	52.3	273	2	Q9PCR3	Q9PCR3 xylella fas
28	45	52.3	412	2	Q916P7	Q916P7 pseudomonas
29	45	52.3	423	2	Q24742	Q24742 bacteroides
30	45	52.3	443	10	Q9S751	Q9S751 oryza sativ
31	45	52.3	448	2	Q9WYR8	Q9WYR8 thermotoga
32	45	52.3	525	10	Q9ATU5	Q9ATU5 lolium rigi
33	45	52.3	525	10	Q9ATU2	Q9ATU2 lolium rigi
34	45	52.3	525	10	Q9ATU1	Q9ATU1 lolium rigi
35	45	52.3	730	10	Q9RG26	Q9RG26 arabidopsis
36	45	52.3	767	12	Q9QUD8	Q9QUD8 tt virus. h
37	45	52.3	1100	11	Q921J9	Q921J9 mus musculu
38	44.5	51.7	766	12	Q91FV0	Q91FV0 tt virus. p
39	44	51.2	143	4	Q9H9A4	Q9H9A4 homo sapien
40	44	51.2	145	2	Q86437	Q86437 pseudomonas
41	44	51.2	257	2	Q56924	Q56924 yersinia en
42	44	51.2	361	4	Q9P1W6	Q9P1W6 homo sapien
43	44	51.2	406	5	Q9W404	Q9W404 drosophila
44	44	51.2	429	5	Q9N8Y2	Q9N8Y2 trypanosoma
45	44	51.2	458	4	Q9UCB1	Q9UCB1 homo sapien

ALIGNMENTS

RESULT 1
ID Q9DUC4 PRELIMINARY: PRT: 723 AA.
AC Q9DUC4;
DT 01-MAR-2001 (TREMBLrel. 16, Created)
DT 01-MAR-2001 (TREMBLrel. 16, Last sequence update)
DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
DE ORF1.
OS TT virus.
OC Viruses; ssDNA viruses; unclassified ssDNA viruses.
OX NCBI_TaxID=68887;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=MF-TTV9;
RA Okamoto H.;
RL Submitted (APR-2000) to the EMBL/Genbank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=MF-TTV9;
RX PubMed=11080484;
RA Okamoto H., Nishizawa T., Tawara A., Peng Y., Takahashi M.,
RA Kishimoto J., Tanaka T., Miyakawa Y., Mayumi M.;
RT "Species-specific TT viruses in humans and nonhuman primates and their
RT phylogenetic relatedness";
RL Virology 277:368-378(2000).
DR EMBL: AB041959; BAB19313.1;
DR InterPro: IPR001563; Serine-carboxypept.
DR PROSITE: PS00131; CARBOXYPEPT_SER_SER; UNKNOWN.1.
SQ SEQUENCE 723 AA; 85393 MW; 232D003098766344 CRC64;

Query Match 66.3%; Score 57; DB 12; Length 723;
Best Local Similarity 100.0%; Pred. No. 4.0;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 5 PMWPMWR 11
Db 2 PMWPMWR 8
RESULT 2
Q9JH31

ID 09JH31 PRELIMINARY; PRT; 746 AA.
 AC 09JH31;
 DT 01-OCT-2000 (TREMBlrel. 15, Created)
 DT 01-OCT-2000 (TREMBlrel. 15, Last sequence update)
 DE 01-OCT-2000 (TREMBlrel. 15, Last annotation update)
 DE ORF1.
 OS TT virus.
 OC Viruses; ssDNA viruses; unclassified ssDNA viruses.
 NC NCB1_TaxID=68887;
 RN 11;
 RP SEQUENCE FROM N.A.
 RC STRAIN=TUN02;
 RA Okamoto H.;
 RL Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
 RN 12;
 RP SEQUENCE FROM N.A.
 RC STRAIN=TUN02;
 RA Ukita M., Okamoto H., Nishizawa T., Tawara A., Takahashi M.,
 RA Iizuka H., Miyakawa Y., Mayumi M.;
 RT "The entire nucleotide sequences of two distinct TT virus (TTV)
 RT isolates (TUN01 and TUN02) remotely related to the original TTV
 RT isolates.";
 RL Arch. Virol. 0:0-0(2000).
 DR EMBL: AB028669; BAA94878.1;
 SQ SEQUENCE 746 AA; 88561 MW; E0B22953AE764E3E CRC64;

Query Match 61.6%; Score 53; DB 12; Length 746;
 Best Local Similarity 54.5%; Pred. No. 15;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 2 LRPMWMPW 12
 Db 1 MANGWRRRR 11

RESULT 3
 0990M4 PRELIMINARY; PRT; 1173 AA.
 AC 0990M4;
 DT 01-JUN-2001 (TREMBlrel. 17, Created)
 DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
 DE 01-JUN-2001 (TREMBlrel. 17, Last annotation update)
 DE SPIKE GLYCOPROTEIN.
 GN S.
 OS Human coronavirus (strain 229E).
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
 OC Coronaviridae; Coronavirus.
 NC NCB1_TaxID=11137;
 RN 11;
 RP SEQUENCE FROM N.A.
 RC STRAIN=229E;
 RA Bonavia A., Holmes K.V.;
 RT "Viral and cellular changes in a human cell line persistently infected
 RT with human coronavirus HCoV-229E.";
 RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF344186; AAK32188.1;
 SQ SEQUENCE 1173 AA; 128669 MW; ABC6E0A75E8BD8A4 CRC64;

Query Match 61.6%; Score 53; DB 12; Length 1173;
 Best Local Similarity 62.5%; Pred. No. 23;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 LRPMWMPW 9
 Db 1112 IKPMWVW 1119

RESULT 4
 0990M3 PRELIMINARY; PRT; 1173 AA.
 AC 0990M3;

DT 01-JUN-2001 (TREMBlrel. 17, Created)
 DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
 DE 01-JUN-2001 (TREMBlrel. 17, Last annotation update)
 DE SPIKE GLYCOPROTEIN.
 GN S.
 OS Human coronavirus (strain 229E).
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
 NC NCB1_TaxID=11137;
 RN 11;
 RP SEQUENCE FROM N.A.
 RC STRAIN=229E;
 RA Bonavia A., Holmes K.V.;
 RT "Viral and cellular changes in a human cell line persistently infected
 RT with human coronavirus HCoV-229E.";
 RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF344187; AAK32189.1;
 SQ SEQUENCE 1173 AA; 128683 MW; 9E2368160082A81A CRC64;

Query Match 61.6%; Score 53; DB 12; Length 1173;
 Best Local Similarity 62.5%; Pred. No. 23;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 LRPMWMPW 9
 Db 1112 IKPMWVW 1119

RESULT 5
 0990M2 PRELIMINARY; PRT; 1173 AA.
 AC 0990M2;
 DT 01-JUN-2001 (TREMBlrel. 17, Created)
 DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
 DE 01-JUN-2001 (TREMBlrel. 17, Last annotation update)
 DE SPIKE GLYCOPROTEIN.
 GN S.
 OS Human coronavirus (strain 229E).
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
 OC Coronaviridae; Coronavirus.
 NC NCB1_TaxID=11137;
 RN 11;
 RP SEQUENCE FROM N.A.
 RC STRAIN=229E;
 RA Bonavia A., Holmes K.V.;
 RT "Viral and cellular changes in a human cell line persistently infected
 RT with human coronavirus HCoV-229E.";
 RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF344188; AAK32190.1;
 SQ SEQUENCE 1173 AA; 128653 MW; 8B658FCBBD1842DA CRC64;

Query Match 61.6%; Score 53; DB 12; Length 1173;
 Best Local Similarity 62.5%; Pred. No. 23;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 LRPMWMPW 9
 Db 1112 IKPMWVW 1119

RESULT 6
 0990M1 PRELIMINARY; PRT; 1173 AA.
 AC 0990M1;
 DT 01-JUN-2001 (TREMBlrel. 17, Created)
 DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
 DE 01-JUN-2001 (TREMBlrel. 17, Last annotation update)
 DE SPIKE GLYCOPROTEIN.
 GN S.
 OS Human coronavirus (strain 229E).
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;

OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=1137;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=229E;
RA Bonavia A., Holmes K.V.;
RT "Viral and cellular changes in a human cell line persistently infected
with human coronavirus HCoV-229E."
RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF344189; AAK32191.1; -
SQ SEQUENCE 1173 AA; 128760 MW; B73A165A6270152A CRC64;

Query Match
Best Local Similarity 61.6%; Score 53; DB 12; Length 1173;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 LRMPMPW 9
Db 1112 IKMPMPW 1119

RESULT 7
ID 084712 PRELIMINARY; PRT; 1383 AA.
AC 084712;
DT 01-NOV-1996 (TREMBlrel. 01, Created)
DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
DT 01-JUN-2001 (TREMBlrel. 17, Last annotation update)
DE SPIKE PROTEIN.
OS porcine epidemic diarrhea virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=28295;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BRI/87;
RX MEDLINE=94231173; PubMed=8176382;
RA Duarte M., Laude H.;
RT "Sequence of the spike protein of the porcine epidemic diarrhoea
virus."
RL J. Gen. Virol. 75:1195-1200(1994).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=BRI/87;
RX MEDLINE=93389433; PubMed=8397280;
RA Bridgen A., Duarte M., Tobler K., Laude H., Ackermann M.;
RT "Sequence determination of the nucleocapsid protein gene of the
porcine epidemic diarrhoea virus confirms that this virus is a
coronavirus related to human coronavirus 229E and porcine
transmissible gastroenteritis virus."
RL J. Gen. Virol. 74:1795-1804(1993).
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=BRI/87;
RX MEDLINE=94120721; PubMed=8291230;
RA Duarte M., Tobler K., Bridgen A., Rasschaert D., Ackermann M.,
Laude H.;
RT "Sequence analysis of the porcine epidemic diarrhoea virus genome
between the nucleocapsid and spike protein genes reveals a polymorphic
ORF."
RL Virology 198:466-476(1994).
DR EMBL: Z25483; CAA80971.1; -
DR InterPro: IPR002551; Corona_S1.
DR InterPro: IPR002552; Corona_S2.
DR Pfam: PF01600; Corona_S1; 1.
DR Pfam: PF01601; Corona_S2; 1.
FT CONFLICT 422 422 Y -> N (IN REF. 1).
SQ SEQUENCE 1383 AA; 151404 MW; 741C84D5DD3BDC4D CRC64;

Query Match 61.6%; Score 53; DB 12; Length 1383;
Best Local Similarity 62.5%; Pred. No. 27;

Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
OY 2 LRMPMPW 9
Db 1321 IKMPMPW 1328

RESULT 8
ID 0977V5 PRELIMINARY; PRT; 1245 AA.
AC 0977V5;
DT 01-NOV-1999 (TREMBlrel. 12, Created)
DT 01-NOV-1999 (TREMBlrel. 12, Last sequence update)
DT 01-NOV-1999 (TREMBlrel. 12, Last annotation update)
DE CONIDIOSPORE SURFACE PROTEIN.
GN CMPI.
OS Trichoderma harzianum.
OC Eukaryota; Fungi; Ascomycota; mitosporic Ascomycota; Trichoderma.
OX NCBI_TaxID=5544;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 32173;
RA Puyesky M., Benhamou N., Ponce Noyola P., Bauw G., Ziv T.,
van Montagu M., Herrera Estrella A., Horwitz B.A.;
RT "Developmental regulation of a gene encoding a multidomain
RT conidiospore surface protein of Trichoderma, cmpl."
RL Submitted (MAR-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL: AJ133651; CAB40845.1; -
SQ SEQUENCE 1245 AA; 135824 MW; 3249C749AFA0CDF8 CRC64;

Query Match 60.5%; Score 52; DB 3; Length 1245;
Best Local Similarity 60.0%; Pred. No. 33;
Matches 6; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

OY 3 RMPMPMPRRK 12
Db 1185 RMQWMSMPRR 1194

RESULT 9
ID 09Y4N1 PRELIMINARY; PRT; 299 AA.
AC 09Y4N1;
DT 01-NOV-1999 (TREMBlrel. 12, Created)
DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)
DT 01-MAY-2000 (TREMBlrel. 13, Last annotation update)
DE HYPOTHETICAL 34.0 KDA PROTEIN (FRAGMENT).
GN DKFZP434C192.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=TESTIS;
RA Ansoorge W., Winkler U., Mewes H.W., Gassenhuber J., Wiemann S.;
RL Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL: AL096753; CAB46428.2; -
KW Hypothetical protein.
FT NON_TER 1
SQ SEQUENCE 299 AA; 34032 MW; 6B8DB606EA88239A CRC64;

Query Match 59.3%; Score 51; DB 4; Length 299;
Best Local Similarity 85.7%; Pred. No. 12;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 5 PMMPMPRR 11
Db 37 PMMPMPRR 43

```
RESULT 10
P96143
ID P96143 PRELIMINARY PRT: 504 AA.
AC P96143:
DT 01-MAY-1997 (TRENBLREL. 03, Created).
DT 01-MAY-1997 (TRENBLREL. 03, Last sequence update).
DT 01-JUN-2001 (TRENBLREL. 17, Last annotation update).
DE PEPTIDE HYDROLASE.
GN TPEL.
OS Thermoactinomyces vulgaris.
OC Bacteria; Firmicutes; Bacillus/Clostridium group; Thermoactinomyces.
NCBI_TaxID=2026;
RN [1]
RP SEQUENCE OF 1-431 FROM N.A.
RC STRAIN=94-2A;
RA Hofmeister J.W.;
RL Thesis (1995), Molecular Genetics,
  Institut fuer Pflanzen-genetik und Kulturpflanzenforschung.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=94-2A;
RA Hofmeister J.W.;
RL Submitted (DEC-1996) to the EMBL/GenBank/DBJ databases.
DR EMBL; 283218; CAB05671.1; -.
DR HSSP; P00800; 1HYT.
DR Interpro; IPR001570; Peptidase_M4.
DR Pfam; PF01447; Peptidase_M4; 1.
DR Hydrolase.
SQ SEQUENCE 504 AA; 56653 MW; 5A7BC05C5AD1315 CRC64;

Query Match
Best Local Similarity 59.3%; Score 51; DB 2; Length 504;
Pred. No. 20;
Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Oy 1 ILRPMWPMR 10
Db 71 LVKMTWPMR 80

RESULT 11
Q18753
ID Q18753 PRELIMINARY PRT: 111 AA.
AC Q18753:
DT 01-NOV-1996 (TRENBLREL. 01, Created).
DT 01-NOV-1996 (TRENBLREL. 01, Last sequence update).
DT 01-NOV-1998 (TRENBLREL. 08, Last annotation update).
DE GLYCINE-RICH.
GN C50P7.8.
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidae;
  Rhabditidae; Peleoderinae; Caenorhabditis.
NCBI_TaxID=6239;
RN [1]
RP SEQUENCE FROM N.A.
RC MEDLINE=94150718; PubMed=7906398;
RA Wilson R., Almscough R., Anderson K., Baynes C., Berks M.,
  Bonfield J., Burton J., Connell M., Copsey T., Cooper J., Coulson A.,
  Craxton M., Dear S., Du Z., Durbin R., Favello A., Fulton L.,
  Gardner A., Green P., Hawkins T., Hillier L., Jier M., Johnston L.,
  Jones M., Kershaw J., Kirsten J., Laister N., Latreille P.,
  Lightning J., Lloyd C., Murray A., Mortimore B., O'Callaghan M.,
  Parsons J., Percy C., Rifkin L., Roopra A., Saunders D., Shownkeen R.,
  Smaldon N., Smith A., Sonhammer E., Staden R., Sulston J.,
  Thierly-Mieg J., Thomas K., Vaubin M., Vaughan K., Waterston R.,
  Watson A., Weinstock L., Wilkinson-Sproat J., Wohlman P.;
  "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
  elegans.";
RT Nature 368:32-38(1994).
RN [2]
RP SEQUENCE FROM N.A.
RA Johnson D., Steilys L.;
RL Submitted (DEC-1995) to the EMBL/GenBank/DBJ databases.
```

```
RN [3]
RP SEQUENCE FROM N.A.
RA Waterston R.;
RL Submitted (NOV-1995) to the EMBL/GenBank/DBJ databases.
DR EMBL; U41557; AAA8303.1; -.
SQ SEQUENCE 111 AA; 10139 MW; 6E729A2E0F9762B9 CRC64;

Query Match
Best Local Similarity 58.1%; Score 50; DB 5; Length 111;
Pred. No. 7;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

Oy 1 ILRPMWPMR 11
Db 12 VWRPMWPMGR 22

RESULT 12
Q9CZAL
ID Q9CZAL PRELIMINARY PRT: 141 AA.
AC Q9CZAL:
DT 01-JUN-2001 (TRENBLREL. 17, Created).
DT 01-JUN-2001 (TRENBLREL. 17, Last sequence update).
DT 01-JUN-2001 (TRENBLREL. 17, Last annotation update).
DE 2810031J10RIK PROTEIN.
GN 2810031J10RIK.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=EMBRYO;
RX MEDLINE=21085660; PubMed=11217851;
RA Kawai J., Shingawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
  Aikawa T., Hara A., Fukunishi Y., Konno H., Adachi S., Fukuda S.,
  Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yananaka I.,
  Saito T., Okazaki Y., Gojobori T., Bono H., Kasukawa T., Saito R.,
  Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
  Fleischmann W., Gaasterland T., Gissi C., King B., Kochia H.,
  Kuehl P., Lewis S., Matsuo Y., Nikaide I., Pesole G., Quackenbush J.,
  Schriml L.M., Staudt F., Suzuki R., Tomita M., Wagner L., Washio T.,
  Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
  Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
  Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
  Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,
  Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
  Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
  Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
  Suzuki H., Toyooka K., Wang K.H., Weitz C., Whitlaker C., Wilming L.,
  Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohlsuki S.,
  Hayashizaki Y.;
  "Functional annotation of a full-length mouse cDNA collection.";
RT Nature 409:685-690(2001).
RN [2]
RP EMBL; AK012846; BAB28508.1; -.
DR MGI; MGI:1919917; 2810031J10RIK.
DR Interpro; IPR003309; SCAN.
DR Pfam; PF02023; SCAN; 1.
DR SMART; SM00431; LER; 1.
DR PROSITE; PS50804; SCAN_BOX; 1.
SQ SEQUENCE 141 AA; 15993 MW; 865C6B735BF8203D CRC64;

Query Match
Best Local Similarity 58.1%; Score 50; DB 11; Length 141;
Pred. No. 8.7;
Matches 6; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

Oy 1 ILRPMWPMR 9
Db 105 VSRPMWPMR 113

RESULT 13
```

```

Q9AUN3
ID Q9AUN3 PRELIMINARY; PRT: 327 AA.
AC Q9AUN3:
DT 01-JUN-2001 (TREMBLrel. 17, Created)
DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
DE 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
DE HYPOTHETICAL PROTEIN.
OS Oryza sativa (Rice).
OC Eukaryota: Viridiplantae: Streptophyta: Embryophyta: Tracheophyta:
OC Spermatophyta: Magnoliophyta: Liliopsida: Poales; Poaceae;
OC Ehrhartoideae; Oryzeae; Oryza.
OX NCBI_TaxID=4530;
RN [1]
RP SEQUENCE FROM N.A.
RA Spiegel L.A., King L., Kirchoff K.A., de la Bastide M., Preston R.R.,
RA Nascimento L.U., Vil M.D., Baker J.P., Miller B., Cunnis D.M.,
RA Kult K.H., Rodriguez S., Santos L., Zutavern T., Ballja V.S.,
RA Shad R.S., Bahret A., Bal H.P., O'Shaughnessy A., Dedhia N.N.,
RA McCombie W.R.;
RT "Genomic Sequence For Oryza sativa, Nipponbare Strain, Chromosome X,
RT Clone OSJNBa0058B19, Complete Sequence.";
RL Submitted (MAR-2001) to the EMBL/Genbank/DBJ databases.
DR EMBL; AC083945; AAK13143.1; -.
SQ SEQUENCE 327 AA; 36672 MW; 5CCA9080664BD0CA CRC64;

Query Match 58.1%; Score 50; DB 10; Length 327;
Best Local Similarity 100.0%; Pred. No. 18;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 6 WMPWRR 11
DB 119 WMPWRR 124

RESULT 14
Q9DUC9 PRELIMINARY; PRT: 735 AA.
AC Q9DUC9:
DT 01-MAR-2001 (TREMBLrel. 16, Created)
DT 01-MAR-2001 (TREMBLrel. 16, Last sequence update)
DT 01-MAR-2001 (TREMBLrel. 16, Last annotation update)
DE ORF1.
OS TT virus.
OC Viruses; ssDNA viruses; unclassified ssDNA viruses.
OX NCBI_TaxID=68887;
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN=PT-TTV6;
RA Okamoto H.;
RL Submitted (Apr-2000) to the EMBL/Genbank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RA STRAIN=PT-TTV6;
RX PubMed=11080484;
RA Okamoto H., Nishizawa T., Tawara A., Peng Y., Takahashi M.,
RA Kishimoto J., Tanaka T., Miyakawa Y., Mayumi M.;
RT "Species-specific TT viruses in humans and nonhuman primates and their
RT phylogenetic relatedness.";
RL Virology 277:368-378(2000).
DR EMBL; AB041957; BAB19308.1; -.
SQ SEQUENCE 735 AA; 86132 MW; 9ED818D6BE6FA5D3 CRC64;

Query Match 58.1%; Score 50; DB 12; Length 735;
Best Local Similarity 46.7%; Pred. No. 37;
Matches 7; Conservative 2; Mismatches 2; Indels 4; Gaps 1;

OY 2 LRPWPM-----WPMRRK 12
DB 1 MAMPWRRRRRRWRRR 15

```

```

RESULT 15
Q9DT80 PRELIMINARY; PRT: 49 AA.
AC Q9DT80:
DT 01-MAR-2001 (TREMBLrel. 16, Created)
DT 01-MAR-2001 (TREMBLrel. 16, Last sequence update)
DT 01-MAR-2001 (TREMBLrel. 16, Last annotation update)
DE 01-MAR-2001 (TREMBLrel. 16, Last annotation update)
DE ORF1 (FRAGMENT).
OS TT virus.
OC Viruses; ssDNA viruses; unclassified ssDNA viruses.
OX NCBI_TaxID=68887;
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN=TYM9;
RX MEDLINE=20568739; PubMed=1118348;
RA Okamoto H., Nishizawa T., Tawara A., Takahashi M., Kishimoto J.,
RA Sai T., Sugai Y.;
RT "TT virus mRNAs detected in the bone marrow cells from an infected
RT individual.";
RL Biochem. Biophys. Res. Commun. 279:700-707(2000).
DR EMBL; AB050449; BAB19930.1; -.
FT NON_TER 49
SQ SEQUENCE 49 AA; 7225 MW; 1DA6F81AB69AA43 CRC64;

Query Match 57.0%; Score 49; DB 12; Length 49;
Best Local Similarity 36.8%; Pred. No. 4.6;
Matches 7; Conservative 2; Mismatches 2; Indels 8; Gaps 1;

OY 2 LRPWPM-----WPMRRK 12
DB 1 MAMPWRRRRRRWRRR 19

```

Search completed: January 4, 2002, 08:47:26
Job time: 414 sec

THIS PAGE BLANK (USPTO)

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:47:48 ; Search time 18.1 Seconds
(without alignments)
24.308 Million cell updates/sec

Title: US-09-444-281-36
Perfect score: 86
Sequence: 1 ILRMPMPWRRK 12

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 100059 seqs, 36664827 residues

Total number of hits satisfying chosen parameters: 100059

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SwissProt_39.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	70	81.4	144	1	INDC_BOVIN
2	53	61.6	1173	1	VGL2_CVB22
3	49	57.0	492	1	ADRO_BOVIN
4	47	54.7	253	1	Y945_MYCTU
5	47	54.7	715	1	YD55_MYCTU
6	45.5	52.9	505	1	TRPE_PSRSS
7	45	52.3	196	1	YA05_SCHPO
8	45	52.3	1108	1	CN38_RAT
9	44	51.2	361	1	FUT3_HUMAN
10	44	51.2	372	1	FUT3_PANTR
11	44	51.2	397	1	MM6_MYCTU
12	44	51.2	535	1	YD6_SCHPO
13	44	51.2	967	1	MM4_MYCTU
14	44	51.2	968	1	MM2_MYCTU
15	44	51.2	984	1	SLX3_MOUSE
16	44	51.2	1154	1	VGL2_IBVD2
17	44	51.2	1162	1	VGL2_IBVK
18	44	51.2	1162	1	VGL2_IBVK
19	44	51.2	1162	1	VGL2_IBVM
20	44	51.2	1163	1	VGL2_IBVM
21	43.5	50.6	276	1	KCEL_RHOPA
22	43.5	50.6	2436	1	ABC2_HUMAN
23	43	50.0	51	1	LHB2_ECTHA
24	43	50.0	711	1	MM4_STRCO
25	43	50.0	958	1	MM4_MYCTU
26	43	50.0	1112	1	CN3B_HUMAN
27	43	50.0	1225	1	VGL2_CVPR8
28	43	50.0	1225	1	VGL2_CVPRM
29	43	50.0	1235	1	VGL2_CVMAH
30	43	50.0	1324	1	VGL2_CVMA5
31	43	50.0	1353	1	VGL2_CVHOC
32	43	50.0	1363	1	VGL2_CVPR
33	43	50.0	1363	1	VGL2_CVBL9

34	43	50.0	1363	1	VGL2_CVBLY	P25192 bovine coro
35	43	50.0	1363	1	VGL2_CVBH	P15777 bovine coro
36	43	50.0	1363	1	VGL2_CVBH	P25193 bovine coro
37	43	50.0	1363	1	VGL2_CVBH	P25194 bovine coro
38	43	50.0	1376	1	VGL2_CVMA	P22432 murine coro
39	43	50.0	1376	1	VGL2_CVMA	P02385 murine coro
40	43	50.0	1447	1	VGL2_CVPR	P02167 porcine tra
41	43	50.0	1447	1	VGL2_CVPR	P07946 porcine tra
42	43	50.0	1447	1	VGL2_CVPR	P01977 porcine tra
43	43	50.0	1449	1	VGL2_CVPR	P18450 porcine tra
44	43	50.0	1449	1	VGL2_CVPR	P33470 porcine tra
45	43	50.0	1451	1	VGL2_CVCAI	P36300 canine ente

ALIGNMENTS

RESULT 1	INDC_BOVIN	STANDARD	PRT	144 AA.
AC	P33046:			
DT	01-OCT-1993 (Rel. 27, Created)			
DT	01-OCT-1993 (Rel. 27, Last sequence update)			
DT	01-NOV-1997 (Rel. 35, Last annotation update)			
DE	INDOLICIDIN PRECURSOR.			
OS	Bos taurus (Bovine).			
OC	Eukaryota: Metazoa: Chordata: Vertebrata: Euteleostomi:			
OC	Mammalia: Eutheria: Cetartiodactyla: Ruminantia: Pecora: Bovidea:			
OC	Bovidae: Bovinae: Bos.			
OX	NCBI_TaxID=9913;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Bone marrow:			
RC	MEDLINE=92392368; PubMed=1520337;			
RA	del Sal G., Stortel P., Schneider C., Romeo D., Zanetti M.;			
RT	"cDNA cloning of the neutrophil bactericidal peptide indolicidin.";			
RL	Biochem. Biophys. Res. Commun. 187:467-472(1992).			
RN	[2]			
RP	SEQUENCE OF 131-143.			
RC	TISSUE=Neutrophils:			
RC	MEDLINE=92165771; PubMed=1537821;			
RA	Seisted M.E., Novotny M.J., Morris W.L., Tang Y.-O., Smith W.;			
RA	"Indolicidin, a novel bactericidal tridecapeptide amide from			
RT	neutrophils.";			
RL	J. Biol. Chem. 267:4292-4295(1992).			
CC	- FUNCTION: POTENT MICROBICIDAL ACTIVITY, ACTIVE AGAINST			
CC	STAPHYLOCOCCUS AUREUS AND ESCHERICHIA COLI.			
CC	- TISSUE SPECIFICITY: LARGE GRANULES OF NEUTROPHILS.			
CC	- PWM: ELASTASE MIGHT BE RESPONSIBLE FOR ITS MATURATION.			
CC	- SIMILARITY: BELONGS TO THE CATHELICIDIN FAMILY.			
CC	-----			
CC	This SWISS-PROT entry is copyright. It is produced through a collaboration			
CC	between the Swiss Institute of Bioinformatics and the EMBL outstation -			
CC	the European Bioinformatics Institute. There are no restrictions on its			
CC	use by non-profit institutions as long as its content is in no way			
CC	modified and this statement is not removed. Usage by and for commercial			
CC	entities requires a license agreement (See http://www.isb-sib.ch/announce/			
CC	or send an email to license@sib-sib.ch).			
CC	-----			
DR	EMBL: X67340; CAA47755.1; -			
DR	PIR: JCI222; JCI222.			
DR	PIR: A42387; A42387.			
DR	InterPro: IPR001894; Cathelicidin.			
DR	Pfam: PF00666; Cathelicidins; 1.			
DR	ProDom: PD001838; Cathelicidin; 1.			
DR	PROSITE: PS00946; CATHELICIDINS_1; 1.			
DR	PROSITE: PS00947; CATHELICIDINS_2; 1.			
FW	Antibiotic: Amidation; Signal.			
KW	SIGNAL			
FT	PROPEP	1	29	POTENTIAL.
FT	PEPTIDE	131	143	INDOLICIDIN, CARBOXYLIC ACID (BY
FT	MOD_RES	30	30	PYRROLIDONE

FT DISULFID 85 96 SIMILARITY.
 FT DISULFID 107 124 BY SIMILARITY.
 FT MOD_RES 143 143 AMIDATION (G-144 PROVIDE AMIDE GROUP).
 SQ SEQUENCE 144 AA: 16479 MW; E3B1CBHE55C09911 CRC64;

Query Match 81.4%; Score 70; DB 1; Length 144;
 Best Local Similarity 88.9%; Pred. No. 0.0039; Indels 0; Gaps 0;
 Matches 8; Conservative 1; Mismatches 0;

OY 3 RWPMPWRR 11
 Db 135 KWPMPWRR 143

RESULT 2
 VGL2_CVH22 STANDARD; PRT: 1173 AA.
 ID VGL2_CVH22
 AC P15423;
 DT 01-APR-1990 (rel. 14, Created)
 DT 01-APR-1990 (rel. 14, Last sequence update)
 DT 15-JUL-1999 (rel. 38, Last annotation update)
 DE E2 GLYCOPROTEIN PRECURSOR (SPIKE GLYCOPROTEIN) (PEPLOMER PROTEIN).
 GN S.
 OS Human coronavirus (strain 229E).
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
 OC Coronaviridae; Coronavirus.
 OX NCBI_TaxID=11137;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=90264837; PubMed=2345367;
 RA Raabe T., Schelle-Prinz B., Siddell S.G.;
 RT "Nucleotide sequence of the gene encoding the spike glycoprotein of
 human coronavirus HCV 229E".
 RL J. Gen. Virol. 71:1065-1073(1990).
 CC -1- FUNCTION: THE PELOMER PROTEIN MEDIATES THE BINDING OF VIRIONS
 TO THE HOST CELL RECEPTOR AND IS INVOLVED IN MEMBRANE FUSION
 AND IN SYNCTYUM FORMATION.
 CC -1- SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 between the Swiss Institute of Bioinformatics and the EMBL outstation -
 the European Bioinformatics Institute. There are no restrictions on its
 use by non-profit institutions as long as its content is in no way
 modified and this statement is not removed. Usage by and for commercial
 entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL: X16816; CAA34723.1; -
 DR PIR: A34766; VGIIHC.
 DR InterPro: IPR002551; Corona_S1.
 DR InterPro: IPR002552; Corona_S2.
 DR Pfam: PF01600; Corona_S1; 1.
 DR Pfam: PF01601; Corona_S2; 1.
 KW Glycoprotein; Envelope protein; Transmembrane; Signal.
 FT SIGNAL 1 15
 FT CHAIN 16 1173
 FT DOMAIN 16 1115
 FT TRANSMEM 1116 1135
 FT DOMAIN 1136 1173
 FT DOMAIN 1136 1157
 FT CARBOHYD 23 23
 FT CARBOHYD 62 62
 FT CARBOHYD 98 98
 FT CARBOHYD 147 147
 FT CARBOHYD 171 171
 FT CARBOHYD 176 176
 FT CARBOHYD 220 220
 FT CARBOHYD 243 243
 FT CARBOHYD 326 326
 FT CARBOHYD 333 333
 FT CARBOHYD 440 440

FT CARBOHYD 464 464 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 518 518 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 538 538 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 542 542 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 568 568 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 581 581 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 587 587 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 663 663 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 671 671 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 671 671 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 930 930 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1015 1015 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1020 1020 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1037 1037 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1049 1049 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1061 1061 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1066 1066 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1076 1076 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1082 1082 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1096 1096 N-LINKED (GLCNAC. . .) (POTENTIAL).
 SQ SEQUENCE 1173 AA: 128639 MW; B9CA9A41A796B3BD CRC64;

Query Match 61.6%; Score 53; DB 1; Length 1173;
 Best Local Similarity 62.5%; Pred. No. 4.4; Indels 0; Gaps 0;
 Matches 5; Conservative 2; Mismatches 1;

OY 2 LRWPMPW 9
 Db 1112 IKWPMPWV 1119

RESULT 3
 ID ADRO_BOVIN STANDARD; PRT: 492 AA.
 AC P08165;
 DT 01-AUG-1988 (rel. 08, Created)
 DT 15-JUL-1998 (rel. 36, Last sequence update)
 DT 20-AUG-2001 (rel. 40, Last annotation update)
 DE NADP:ADRENODOXIN OXIDOREDUCTASE, MITOCHONDRIAL PRECURSOR
 DE (EC 1.18.1.2) (ADRENODOXIN REDUCTASE) (AR) (FERREDOXIN-NADP(+)
 DE REDUCTASE).
 GN FDXR OR ADXR.
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovidae; Bovinae; Bos.
 OX NCBI_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A., AND ALTERNATIVE SPLICING.
 RX MEDLINE=94177140; PubMed=8130767;
 RA Takata Y., Sagara Y., Kono A., Sekimizu K., Horiuchi T.;
 RT "Gene structure of bovine adrenodoxin reductase.";
 RL Biol. Pharm. Bull. 16:1200-1206(1993).
 RN [2]
 RP SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.
 RX MEDLINE=88198050; PubMed=3448086;
 RA Sagara Y., Takata Y., Miyata T., Hara T., Horiuchi T.;
 RT "Cloning and sequence analysis of adrenodoxin reductase cDNA from
 bovine adrenal cortex.";
 RL J. Biochem. 102:1333-1336(1987).
 RN [3]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=87270696; PubMed=3038094;
 RA Nonaka Y., Murakami H., Yabusaki Y., Kuramitsu S., Kagamiyama H.,
 Yamano T., Okamoto M.;
 RT "Molecular cloning and sequence analysis of full-length cDNA for mRNA
 of adrenodoxin oxidoreductase from bovine adrenal cortex.";
 RL Biochem. Biophys. Res. Commun. 145:1239-1247(1987).
 RN [4]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Adrenal cortex;
 RX MEDLINE=89170752; PubMed=2924777;
 RA Hanukoglu I., Gutfinger T.;

RT "cDNA sequence of adrenodoxin reductase. Identification of NADP-binding sites in oxidoreductases." RT Eur. J. Biochem. 180:479-484(1989).

RN [5]

RP SEQUENCE OF N-TERMINUS, AND PARTIAL SEQUENCE. RP TISSUE-Adrenal cortex; MEDLINE=88082777; PubMed=3691502; RA Hannukoglu I., Gutfinger T., Hanlu M., Shively J.E.; RT "Isolation of a cDNA for adrenodoxin reductase (ferredoxin-NADP+ reductase). Implications for mitochondrial cytochrome P-450 systems." RT Eur. J. Biochem. 169:449-455(1987).

RN [6]

RP X-RAY CRYSTALLOGRAPHY (1.7 ANGSTROMS) OF 33-492. RP TISSUE-Adrenal gland; MEDLINE=99299392; PubMed=10369776; RA Ziegler G.A., Vonnheim C., Hannukoglu I., Schulz G.E.; RT "The structure of adrenodoxin reductase of mitochondrial P450 systems: electron transfer for steroid biosynthesis." J. Mol. Biol. 289:981-990(1999).

CC -1- FUNCTION: SERVES AS THE FIRST ELECTRON TRANSFER PROTEIN IN ALL THE CC MITOCHONDRIAL P450 SYSTEMS, INCLUDING CHOLESTEROL SIDE CHAIN CC CLEAVAGE IN ALL STEROIDGENIC TISSUES, STEROID 11-BETA CC HYDROXYLATION IN THE ADRENAL CORTEX, 25-OH-VITAMIN D3-24 CC HYDROXYLATION IN THE KIDNEY, AND STEROL C-27 HYDROXYLATION IN THE CC LIVER.

CC -1- CATALYTIC ACTIVITY: REDUCED ADRENODOXIN + NADP(+) = OXIDIZED CC ADRENODOXIN + NADPH.

CC -1- COFACTOR: FAD.

CC -1- PATHWAY: CHOLESTEROL SIDE-CHAIN-CLEAVAGE SYSTEM.

CC -1- SUBUNIT: MONOMER.

CC -1- SUBCELLULAR LOCATION: MITOCHONDRIAL MATRIX.

CC -1- ALTERNATIVE PRODUCTS: 2 ISOFORMS. A SHORT FORM (SHOWN HERE) AND A CC LONG FORM. ARE PRODUCED BY ALTERNATIVE SPLICING. THE LONG FORM CC REPRESENTS 10-20% OF ALL ADRENODOXIN REDUCTASE MRNA. AND SEEMS TO CC BE INACTIVE.

CC -----

CC This SWISS-PROT entry is copyright. It is produced through a collaboration CC between the Swiss Institute of Bioinformatics and the EMBL outstation - CC the European Bioinformatics Institute. There are no restrictions on its CC use by non-profit institutions as long as its content is in no way CC modified and this statement is not removed. Usage by and for commercial CC entities requires a license agreement (see <http://www.isb-sib.ch/announce/> or send an email to license@sib-sib.ch).

CC -----

DR EMBL: M17029; AAA0362.1; -

DR EMBL: D00211; BAA00150.1; -

DR EMBL: X13736; CAA32002.1; -

DR PIR: A29604; A29604.

DR PIR: J50390; J50390.

DR PIR: S03558; S03558.

DR PIR: J70751; J70751.

DR PDB: 1CJC; 12-APR-99.

DR PDB: 1E1L; 02-JUN-00.

DR InterPro: IPR000759; Adnrdx_redctase.

DR PRINTS: PR00419; ADXRDPASE.

DR Electron transport: Oxidoreductase; Flavoprotein; NADP: FAD; KM Mitochondrion; Transit peptide; Alternative splicing; 3D-structure.

FT TRANSIT 1 32 MITOCHONDRION.

FT CHAIN 33 492 NADPH:ADRENODOXIN OXIDOREDUCTASE.

FT VARSPPLIC -204 204 E -> EVLLICQ (IN LONG ISOFORM).

FT CONFLICT 77 77 G -> R (IN REF. 3).

FT CONFLICT 81 94 FGVADPHEVKNVI -> VWLALTTPRSMLL (IN REF. 3).

FT CONFLICT 124 128 QDAYH -> RYRRLT (IN REF. 3).

FT CONFLICT 268 268 K -> R (IN REF. 3).

FT CONFLICT 317 318 PS -> RL (IN REF. 3).

FT CONFLICT 323 333 RAAGIRLAVTR -> ARSASMSPE (IN REF. 3).

FT CONFLICT 341 352 TRAVPTGVEDL -> HFGSAHWGCGP (IN REF. 3).

SO SEQUENCE 492 AA; 54338 MW; E68F6F5D18F53131 CRC64;

Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 4 WRMPWP 9

Db 6 WRMPWP 11

RESULT 4

Y945_MYCTU STANDARD: PRT: 253 AA.

AC P71564;

DT 01-NOV-1997 (Rel. 35, Created)

DT 01-NOV-1997 (Rel. 35, Last sequence update)

DT 20-AUG-2001 (Rel. 40, Last annotation update)

DE PUTATIVE OXIDOREDUCTASE RV0945 (EC 1.-.-.-).

GN RV0945 OR MT0971 OR MYC110D7.29C.

OS Mycobacterium tuberculosis.

OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae; OC Actinomycetales; Corynebacteriaceae; Mycobacteriaceae; Mycobacterium.

OX NCBI_TaxID=1773;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=H37RV;

RX MEDLINE=98295987; PubMed=9634230;

RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D., RA Gordon S.V., Eigmeier K., Gas S., Barry C.E. III, Tekala F., RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R., RA Davies R., Devlin K., Felwell T., Gentles S., Hamlin N., Holroyd S., RA Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy J., RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J., RA Rutter S., Seeger K., Skelton S., Squares R., Squires R., RA Ruiton J.E., Taylor K., Whitehead S., Barrell B.G.; RT "Deciphering the biology of Mycobacterium tuberculosis from the RT complete genome sequence." RT Nature 393:537-544(1998).

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=CDC 1551 / Oshkosh;

RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O., RA Peterson S., Deboy R., Dodson R., Gwinn M.L., Haft D., Hickey E., RA Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L., RA Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A., RA Bishal W.; RT "Whole genome comparison of Mycobacterium tuberculosis clinical and RT laboratory strains." RT Submitted (Apr-2001) to the EMBL/GenBank/DBJ databases.

CC -1- SIMILARITY: BELONGS TO THE SHORT-CHAIN DEHYDROGENASES/REDUCTASES CC (SDR) FAMILY.

CC -----

CC This SWISS-PROT entry is copyright. It is produced through a collaboration CC between the Swiss Institute of Bioinformatics and the EMBL outstation - CC the European Bioinformatics Institute. There are no restrictions on its CC use by non-profit institutions as long as its content is in no way CC modified and this statement is not removed. Usage by and for commercial CC entities requires a license agreement (see <http://www.isb-sib.ch/announce/> or send an email to license@sib-sib.ch).

CC -----

DR EMBL: Z79700; CAB02005.1; -

DR EMBL: AE006982; AAK45219.1; -

DR TIGR: MT0971; -

DR Tuberculist: RV0945; -

DR InterPro: IPR002198; ADH_short.

DR Pfam: PF00106; adh_short.1.

DR PROSITE: PS00061; ADH_SHORT; 1.

DR Hypothetical protein; Oxidoreductase; Complete proteome.

FT ACT SITE 159 159 BY SIMILARITY.

SO SEQUENCE 253 AA; 27138 MW; BAD937208842DA12 CRC64;

Query Match 54.7%; Score 47; DB 1; Length 253;

Best Local Similarity 100.0%; Pred. No. 6.6;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 PMWFW 9
 Db 230 PMWFW 234

RESULT 5
 ID YD55_MYCTU STANDARD; PRT; 715 AA.
 AC 011025;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE HYPOHETICAL 78.2 KDA PROTEIN RV1355C.
 GN RV1355C OR MT1398 OR MTCY02B10.19C.
 OS Mycobacterium tuberculosis.
 OC Bacteria; Firmicutes; Actinobacteriia; Actinobacteridae;
 OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
 NC NCBL_TaxID=1773;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-H37RV;
 RX MEDLINE=98295987; PubMed=9634230;
 RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
 RA Gordon S.V., Eiglmeier K., Gas S., Barry C.E. III, Tekala F.,
 RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
 RA Davies R., Devlin K., Feltwell T., Gentles S., Hamlin N., Holroyd S.,
 RA Hornsby T., Javelin K., Krogh A., McLean J., Moule S., Murphy L.,
 RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
 RA Rutter S., Seeger K., Skelton S., Squares S., Squares R.,
 RA Sulston J.E., Taylor K., Whitehead S., Barrett B.G.,
 RA "Deciphering the biology of Mycobacterium tuberculosis from the
 RT complete genome sequence."
 RL Nature 393:537-544(1998).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CDC 1551 / Oshkosh;
 RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
 RA Peterson J., DeBoy R., Dodson R., Gwin M.L., Hatt D., Hickey E.,
 RA Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L.,
 RA Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A.,
 RA Bishai W.;
 RT "Whole genome comparison of Mycobacterium tuberculosis clinical and
 RL laboratory strains."
 RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
 CC CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL: Z75555; CAA93988.1; -;
 DR EMBL: AE007012; AAK43661.1; ALT_INIT.
 DR TIGR: MT1398; -;
 DR Tuberculist; RV1355C; -;
 DR InterPro: IPR000594; Thif_family.
 DR Pfam: PF00899; Thif_family; 1.
 DR Hypothetical protein, complete proteome.
 KM SEQUENCE 715 AA; 78181 MW; 453495248A56041C CRC64;
 SQ

Query Match 54.7%; Score 47; DB 1; Length 715;
 Best Local Similarity 66.7%; Pred. No. 17;
 Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
 OY 3 RMPWMPWR 11
 Db 65 RMAVYPMWR 73
 RESULT 6

TRPE_PSESS
 ID TRPE_PSESS STANDARD; PRT; 505 AA.
 AC P21689;
 DT 01-MAY-1991 (Rel. 18, Created)
 DT 01-MAY-1991 (Rel. 18, Last sequence update)
 DT 30-MAY-2000 (Rel. 39, Last annotation update)
 DE ANTHRANILATE SYNTHASE COMPONENT I (EC 4.1.3.27).
 GN TRPE.
 OS Pseudomonas syringae (pv. savastanoi).
 OC Bacteria; Proteobacteria; gamma subdivision; Pseudomonadaceae;
 OC Pseudomonas.
 NC NCBL_TaxID=29438;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=91100331; PubMed=1987141;
 RA da Costa E., Silva O., Kosuge T.;
 RT "Molecular characterization and expression analysis of the
 RT anthranilate synthase gene of Pseudomonas syringae subsp.
 RT savastanoi."
 RL J. Bacteriol. 173:463-471(1991).
 CC CC
 CC -I- CATALYTIC ACTIVITY: CHORISMATE + L-GLUTAMINE = ANTHRANILATE +
 CC PYRUVATE + L-GLUTAMATE.
 CC CC
 CC -I- PATHWAY: FIRST STEP IN BIOSYNTHESIS OF TRYPTOPHAN.
 CC CC
 CC -I- SUBUNIT: TETRAMER OF TWO COMPONENTS I AND TWO COMPONENTS II (BY
 CC SIMILARITY).
 CC CC
 CC -I- MISCELLANEOUS: COMPONENT I CATALYZES THE FORMATION OF ANTHRANILATE
 CC USING AMMONIA RATHER THAN GLUTAMINE, WHEREAS COMPONENT II PROVIDES
 CC GLUTAMINE AMIOTRANSFERASE ACTIVITY.
 CC CC
 CC -I- SIMILARITY: BELONGS TO THE ANTHRANILATE SYNTHASE COMPONENT I
 CC FAMILY.
 CC CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL: M55911; AAA26016.1; -;
 DR PIR: A39128; A39128.
 DR InterPro: IPR000350; Chorismate_bind.
 DR Pfam: PF00425; Chorismate_bind; 1.
 DR PRINTS: PR00095; ANTSNTNASEI.
 DR ProDom: PD000779; Chorismate_bind; 1.
 KM Tyrophan biosynthesis; Lyase.
 SQ SEQUENCE 505 AA; 56084 MW; A38E81931331F6BB CRC64;
 SO

Query Match 52.9%; Score 45.5; DB 1; Length 505;
 Best Local Similarity 28.0%; Pred. No. 19;
 Matches 7; Conservative 2; Mismatches 3; Indels 13; Gaps 1;

OY 1 ILRW-----PMWPMWR 12
 Db 467 VLEWERTLNKRRAMVGSAMWPMWR 491
 RESULT 7
 ID YAO5_SCHPO STANDARD; PRT; 196 AA.
 AC Q09677;
 DT 01-NOV-1995 (Rel. 32, Created)
 DT 01-NOV-1995 (Rel. 32, Last sequence update)
 DT 01-NOV-1995 (Rel. 32, Last annotation update)
 DE HYPOHETICAL 22.1 KDA PROTEIN CSH10.05C IN CHROMOSOME I.
 GN SPAC5H10.05C.
 OS Schizosaccharomyces pombe (Fission yeast).
 OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
 OC Schizosaccharomycetales; Schizosaccharomycetaceae;
 OC Schizosaccharomyces.
 NC NCBL_TaxID=4896;
 RN [1]

RP SEQUENCE FROM N.A.
 RC STRAIN-972;
 RA Connor R., Churcher C.M., Barrell B.G., Rajandream M.A., Walsh S.V.;
 RL Submitted (MAY-1995) to the EMBL/GenBank/DBJ databases
 CC -1 SIMILARITY: STRONG, TO BACTERIAL MODULATOR OF DRUG ACTIVITY B (MDAB).
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL: Z49811; CAA89955.1;
 DR InterPro: IPR003680; NADHdh_2;
 DR Pfam: PF02525; NADHdh_2; 1.
 KM Hypothetical protein.
 SQ SEQUENCE 196 AA; 22104 MW; 436764DA9E26074C CRC64;

 Query Match 52.38; Score 45; DB 1; Length 196;
 Best Local Similarity 50.0%; Pred. No. 9.5;
 Matches 8; Conservative 2; Mismatches 2; Indels 4; Gaps 2;

 QY 1 ILRMP-WW---PMRRK 12
 Db 63 IYQWPMWMMCTPMRLK 78

 RESULT 8
 CN3B_RAT STANDARD; PRT: 1108 AA.
 ID AC Q63085;
 DT 15-JUL-1998 (Rel. 36, Created)
 DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DT 15-JUL-1998 (Rel. 36, Last annotation update)
 DE GMP-INHIBITED 3', 5'-CYCLIC PHOSPHODIESTERASE B (EC 3.1.4.17) (CYCLIC
 DE GMP INHIBITED PHOSPHODIESTERASE B) (CGI-PDE B) (CGIPDE1).
 GN PDE3B.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OX NCBI_TaxId=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-SPRAGUE-DAWLEY; TISSUE=adipose tissue;
 RX MEDLINE=93366761; PubMed=8395509;
 RA Taira M., Hockman S.C., Calvo J.C., Taira M., Belfrage P.,
 RA Mangiatello V.C.;
 RT "Molecular cloning of the rat adipocyte hormone-sensitive cyclic GMP-
 RT inhibited cyclic nucleotide phosphodiesterase.";
 RL J. Biol. Chem. 268:18573-18579(1993).
 CC -1- FUNCTION: MAY PLAY A ROLE IN FAT METABOLISM.
 CC -1- CATALYTIC ACTIVITY: GUANOSINE 3', 5'-CYCLIC PHOSPHATE + H(2)O =
 CC GUANOSINE 5'-PHOSPHATE.
 CC -1- ENZYME REGULATION: INHIBITED BY GMP.
 CC -1- SUBCELLULAR LOCATION: MEMBRANE-BOUND (POTENTIAL).
 CC -1- TISSUE SPECIFICITY: ABUNDANT IN ADIPOSE TISSUES.
 CC -1- SIMILARITY: BELONGS TO THE CYCLIC NUCLEOTIDE PHOSPHODIESTERASE
 CC FAMILY.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL: Z22867; CAA80489.1;
 DR InterPro: IPR003607; HDC.

DR InterPro: IPR002073; PDEase.
 DR Pfam: PF00233; PDEase; 1.
 DR SMART: SM00471; HDC; 1.
 DR PROSITE: PS00126; PDEASE_1; 1.
 KM Hydrolase; GMP; Membrane.
 FT DOMAIN 16 22 POLY-PRO.
 FT DOMAIN 99 102 POLY-ALA.
 FT DOMAIN 175 179 POLY-ALA.
 FT DOMAIN 1007 1021 POLY-ASP.
 FT DOMAIN 1068 1071 POLY-GLU.
 FT DOMAIN 1101 1104 POLY-GLU.
 SQ SEQUENCE 1108 AA; 123105 MW; C9B5078C7D3ADD6D CRC64;

 Query Match 52.38; Score 45; DB 1; Length 1108;
 Best Local Similarity 62.5%; Pred. No. 46;
 Matches 5; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

 QY 4 WPMWPRR 11
 Db 164 WQWWSWLR 171

 RESULT 9
 FUT3_HUMAN STANDARD; PRT: 361 AA.
 ID AC P21217; Q99448; Q99449.
 DT 01-MAY-1991 (Rel. 18, Created)
 DT 01-MAY-1991 (Rel. 18, Last sequence update)
 DT 30-MAY-2000 (Rel. 39, Last annotation update)
 DE GALACTOSIDE 3(4)-L-FUCOSYLTRANSFERASE (EC 2.4.1.65) (BLOOD GROUP LEWIS
 DE ALPHA-4-FUCOSYLTRANSFERASE) (LEWIS FT) (FUCOSYLTRANSFERASE 3) (FUCT-
 DE III).
 GN FUT3 OR LE.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 OX NCBI_TaxId=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=91032981; PubMed=1977660;
 RA Kikowska-latallo J.F., Larsen R.D., Nair R.P., Lowe J.B.;
 RT "A cloned human cDNA determines expression of a mouse stage-specific
 RT embryonic antigen and the Lewis blood group
 RT alpha(1,3/1,4)fucosyltransferase.";
 RL Genes Dev. 4:1288-1303(1990).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Liver;
 RX MEDLINE=95378269; PubMed=7650030;
 RA Cameron H.S., Szczepaniak D., Weston M.;
 RT "Expression of human chromosome 19p alpha(1,3)-fucosyltransferase
 RT genes in normal tissues. Alternative splicing, polyadenylation, and
 RT isoforms.";
 RL J. Biol. Chem. 270:20112-20122(1995).
 RN [3]
 RP VARIANT LE(-) MET-105.
 RX MEDLINE=94059067; PubMed=8240322;
 RA Elmgren A., Rydberg L., Larson G.;
 RT "Genotypic heterogeneity among Lewis negative individuals.";
 RL Biochem. Biophys. Res. Commun. 196:515-520(1993).
 RN [4]
 RP VARIANTS LE(-) ARG-20; SER-170 AND ALA-336.
 RX MEDLINE=94059082; PubMed=8240337;
 RA Nishihara S., Iazawa S., Iwasaki H., Nakazato M., Kudo T., Ando T.,
 RA Nishihara H.;
 RT "Alpha (1,3/1,4)fucosyltransferase (FUCT-III) gene is inactivated by
 RT a single amino acid substitution in Lewis histo-blood type negative
 RT individuals.";
 RL Biochem. Biophys. Res. Commun. 196:624-631(1993).
 RN [5]
 RP VARIANTS LE(-) ARG-20 AND SER-170.
 RX MEDLINE=94033579; PubMed=8219240;

RA Koda Y., Kimura H., Mekada E.:
 RT "Analysis of Lewis fucosyltransferase genes from the human gastric
 RT mucosa of Lewis-positive and -negative individuals.";
 RL Blood 82:2915-2919(1993).
 RN [6]
 RP VARIANTS LE(-) ARG-20 AND LYS-356.
 RX MEDLINE=94342259; PubMed=8063716;
 RA Mollitone R., Reguigne I., Kelly R.J., Fletcher A., Watt J.,
 RA Chatfield S., Aziz A., Cameron H.S., Weston B.W., Lowe J.B., Oriol R.:
 RT "Molecular basis for Lewis alpha(1,3/1,4)-fucosyltransferase gene
 RT deficiency (FUT3) found in Lewis-negative Indonesian pedigrees.";
 RL J. Biol. Chem. 269:20987-20994(1994).
 RN [7]
 RP VARIANT LE(-) LYS-356.
 RX MEDLINE=95050753; PubMed=7961897;
 RA Nishihara S., Narimatsu H., Iwasaki H., Yazawa S., Akamatsu S.,
 RA Ando T., Seno T., Narimatsu I.:
 RT "Molecular genetic analysis of the human Lewis histo-blood group
 RT system.";
 RL J. Biol. Chem. 269:29271-29278(1994).
 RN [8]
 RP VARIANTS LE(-) ARG-20; ARG-68; MET-105 AND LYS-356.
 RX MEDLINE=96243526; PubMed=8801770;
 RA Elmgren A., Boerjeson C., Svensson L., Rydberg L., Larson G.:
 RT "DNA sequencing and screening for point mutations in the human Lewis
 RT 'FUT3' gene enables molecular genotyping of the human Lewis blood
 RT group system.";
 RL Vox Sang. 70:97-103(1996).
 RN [9]
 RP VARIANTS LE(-) ARG-68 AND MET-105.
 RX MEDLINE=97413801; PubMed=9268337;
 RA Elmgren A., Mollitone R., Costache M., Boerjeson C., Oriol R.,
 RA Harrington J., Larson G.:
 RT "Significance of individual point mutations, T202C and C314T, in the
 RT human Lewis 'FUT3' gene for expression of Lewis antigens by the human
 RT alpha 1,3/1,4-fucosyltransferase, Fuc-TIII.";
 RL J. Biol. Chem. 272:21994-21998(1997).
 RN [10]
 RP VARIANTS LE(+) K-102; A-124, AND VARIANTS LE(-) N-162; R-223; M-270.
 RX MEDLINE=98366989; PubMed=9703429;
 RA Pang H., Liu Y., Kodai Y., Soejima M., Jia J., Schlaphoff T.,
 RA du Toit E.D., Kimura H.:
 RT "Five novel missense mutations of the Lewis gene 'FUT3' in African
 RT 'Xhosa' and Caucasian populations in South Africa.";
 RL Hum. Genet. 102:675-680(1998).
 CC -I- FUNCTION: MAY CATALYZE ALPHA-1,3 AND ALPHA-1,4 GLYCOSIDIC LINKAGES
 CC INVOLVED IN THE EXPRESSION OF VIM-2, LEWIS A, LEWIS B, STAYL
 CC LEWIS X AND LEWIS X/SSA-1 ANTIGENS. MAY BE INVOLVED IN BLOOD
 CC GROUP LEWIS DETERMINATION; LEWIS-POSITIVE (LE(+)) INDIVIDUALS
 CC HAVE AN ACTIVE ENZYME WHILE LEWIS-NEGATIVE (LE(-)) INDIVIDUALS
 CC HAVE AN INACTIVE ENZYME.
 CC -I- CATALYTIC ACTIVITY: GDP-L-FUCOSE + 1,3-BETA-D-GALACTOSYL-
 CC N-ACETYL-D-GLUCOSAMINYL-R = GDP + 1,3-BETA-D-GALACTOSYL-
 CC (ALPHA-1,4-L-FUCOSYL)-N-ACETYL-D-GLUCOSAMINYL-R.
 CC -I- PATHWAY: GLYCOSYLATION.
 CC -I- SUBCELLULAR LOCATION: TYPE II MEMBRANE PROTEIN. MEMBRANE-BOUND
 CC FORM IN TRANS CISTERNAE OF GOLGI.
 CC -I- TISSUE SPECIFICITY: HIGHLY EXPRESSED IN STOMACH, COLON, SMALL
 CC INTESTINE, LUNG AND KIDNEY AND TO A LESSER EXTENT IN SALIVARY
 CC GLAND, BLADDER, UTERUS AND LIVER.
 CC -I- MISCELLANEOUS: ALSO ACTS ON THE CORRESPONDING 1,4-GALACTOSYL
 CC DERIVATIVE, FORMING 1,3-L-FUCOSYL LINKS.
 CC -I- SIMILARITY: STRUCTURAL SIMILARITY WITH THE OTHER MAMMALIAN
 CC GLYCOSYLTRANSFERASES.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----

DR EMBL: X53578; CAA37641.1; -
 DR EMBL: U27328; AAC50187.1; -
 DR EMBL: U27326; AAC50185.1; -
 DR EMBL: U27327; AAC50186.1; -
 DR EMBL: D89324; BAA13941.1; -
 DR EMBL: D89325; BAA13942.1; -
 DR PIR: A36669; A36669.
 DR MIM: 111100; -
 DR InterPro: IPR001503; Glyco_transf_10.
 DR Pfam: PF00852; Glyco_transf_10; 1..10.
 KW Transferase; Glycosyltransferase; Glycoprotein; Transmembrane;
 KW Signal-anchor; Golgi stack; Polymorphism; Blood group antigen.
 KW Domain
 FT TRANSMEM 16 34
 FT DOMAIN 35 361
 FT CARBOHYD 154 154
 FT CARBOHYD 185 185
 FT VARIANT 20 20
 FT VARIANT 68 68
 FT VARIANT 102 102
 FT VARIANT 105 105
 FT VARIANT 124 124
 FT VARIANT 162 162
 FT VARIANT 170 170
 FT VARIANT 223 223
 FT VARIANT 270 270
 FT VARIANT 336 336
 FT VARIANT 356 356
 FT SEQUENCE 361 AA; 42117 MW; Bf4398044F19C284 CRC64;
 SQ
 Query Match 51.2%; Score 44; DB 1; Length 361;
 Best Local Similarity 85.7%; Pred. No. 23;
 Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 5 PWWPWR 11
 DB 9 PWWPWR 15
 DE
 RESULT 10
 ID FUT3_PANTR STANDARD; PRT; 372 AA.
 AC 019058;
 DT 30-MAY-2000 (Rel. 39, Created)
 DT 30-MAY-2000 (Rel. 39, Last sequence update)
 DT 30-MAY-2000 (Rel. 39, Last annotation update)
 DE GALACTOSIDE 3(4)-L-FUCOSYLTRANSFERASE (EC 2.4.1.65) (BLOOD GROUP LEWIS
 DE ALPHA-4-FUCOSYLTRANSFERASE) (LEWIS FT) (FUCOSYLTRANSFERASE 3) (FUCT-
 DE IIL) (ALPHA-3/4-FUCOSYLTRANSFERASE).
 GN FUT3.
 OS Pan troglodytes (Chimpanzee).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Pan.
 OX NCBI_TaxID=9598;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=98037800; PubMed=9368041;
 RA Costache M., Apoll P.-A., Cailliau A., Elmgren A., Larson G.,
 RA Henry S., Blancher A., Iordachescu D., Oriol R., Mollitone R.:
 RT "Evolution of fucosyltransferase genes in vertebrates.";

RL J. Biol. Chem. 272:29721-29728(1997).
 CC -1- FUNCTION: MAY CATALYZE ALPHA-1,3 AND ALPHA-1,4 GLYCOSIDIC LINKAGES
 CC INVOLVED IN THE EXPRESSION OF STAIYL LEMIS X AND LEMIS X/SEEA-1
 CC ANTIGENS. IT MAY BE INVOLVED IN BLOOD GROUP LEMIS DETERMINATION
 CC (BY SIMILARITY).
 CC -1- CATALYTIC ACTIVITY: GDP-L-FUCOSE + 1,3-BETA-D-GALACTOSYL-
 CC N-ACETYL-D-GLUCOSAMINYL-R = GDP + 1,3-BETA-D-GALACTOSYL-
 CC (ALPHA-1,4-L-FUCOSYL)-N-ACETYL-D-GLUCOSAMINYL-R.
 CC -1- PATHWAY: GLYCOSYLATION.
 CC -1- SUBCELLULAR LOCATION: TYPE II MEMBRANE PROTEIN. MEMBRANE-BOUND
 CC FORM IN TRANS CISTERNA OF GOLGI (BY SIMILARITY).
 CC -1- POLYMORPHISM: THERE ARE TWO ALLELES (A AND B). ALLELE A HAS ARG-
 CC 162 AND VAL-304. ALLELE B HAS GLY-162 AND MET-304.
 CC -1- MISCELLANEOUS: ALSO ACTS ON THE CORRESPONDING 1,4-GALACTOSYL
 CC DERIVATIVE, FORMING 1,3-L-FUCOSYL LINKS.
 CC -1- SIMILARITY: STRUCTURAL SIMILARITY WITH THE OTHER MAMMALIAN
 CC GLYCOSYLTRANSFERASES.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL: Y14033; CAA74360.1; -
 DR InterPro: IPR001503; Glyco_transf_10.
 DR Pfam: PF00852; Glyco_transf_10; 1.
 KW transferase; Glycosyltransferase; Glycoprotein; Transmembrane;
 KW Signal-anchor; Golgi stack; Polymorphism.
 FT DOMAIN 1 14 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 15 34 SIGNAL-ANCHOR (TYPE-II MEMBRANE PROTEIN)
 FT (POTENTIAL).
 FT LUMENAL, CATALYTIC (POTENTIAL)...
 FT CARBOHYD 165 372 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 196 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT VARIANT 162 162 R -> G (IN ALLELE B).
 FT VARIANT 304 304 V -> M (IN ALLELE B).
 FT SEQUENCE 372 AA; 43233 MW; 649CBFB8CA7BD74C CRC64;

Query Match 51.2%; Score 44; DB 1; Length 372;
 Best Local Similarity 85.7%; Pred. No. 23;
 Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 5 RWPMPWR 11
 1 11111
 Db 9 RWPMPWR 15

RESULT 11
 MML6_MYCTU STANDARD; PRT; 397 AA.
 ID MML6_MYCTU
 AC Q10773;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last annotation update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE PUTATIVE MEMBRANE PROTEIN MML6.
 GN MML6 OR RV157 OR MT1608 OR MTCY48.08C.
 OS Mycobacterium tuberculosis.
 CC Bacteria; Filicutes; Actinobacteria; Actinobacteridae;
 CC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
 OX NCBI_TaxID=1773;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=H37RV;
 RX MEDLINE=98295987; PubMed=9634230;
 RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
 RA Gordon S.V., Eigemeier K., Gas S., Barry C.E. III, Tekala F.,
 RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
 RA Davies R., Devlin K., Fellwell T., Gentles S., Hamlin N., Holroyd S.,
 RA Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,

RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
 RA Rutter S., Seeger K., Skellon S., Squares S., Squares R.,
 RA Sulten J.E., Taylor K., Whitehead S., Barrell B.G.;
 RT "Deciphering the biology of Mycobacterium tuberculosis from the
 RT complete genome sequence."
 RL Nature 393:537-544(1998).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CDC 1551 / Oshkosh;
 RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
 RA Peterson J., Deboy R., Dodson R., Gwin M.L., Hatt D., Hickey E.,
 RA Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L.,
 RA Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A.,
 RA Bishai W.;
 RT "Whole genome comparison of Mycobacterium tuberculosis clinical and
 RT laboratory strains."
 RL Submitted (APR-2001) to the EMBL/Genbank/DBJ databases.
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
 CC -1- SIMILARITY: BELONGS TO THE MML6 FAMILY.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL: Z74020; CAA98334.1; -
 DR EMBL: AE007027; AAK45875.1; -
 DR TIGR: MT1608; -
 DR Tuberculist; RV1557; -
 KW Hypothetical protein; Transmembrane; Complete proteome.
 FT TRANSMEM 161 181 POTENTIAL.
 FT TRANSMEM 190 210 POTENTIAL.
 FT TRANSMEM 214 262 POTENTIAL.
 FT TRANSMEM 242 262 POTENTIAL.
 FT TRANSMEM 293 313 POTENTIAL.
 FT TRANSMEM 330 350 POTENTIAL.
 FT SEQUENCE 397 AA; 42421 MW; 678DC6E2A472BF4 CRC64;

Query Match 51.2%; Score 44; DB 1; Length 397;
 Best Local Similarity 75.0%; Pred. No. 25;
 Matches 6; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 3 RWPMPWR 10
 1 11111
 Db 351 RWPMPWR 358

RESULT 12
 YDW6_SCHPO STANDARD; PRT; 535 AA.
 ID YDW6_SCHPO
 AC O13912;
 DT 15-JUL-1998 (Rel. 36, Created)
 DT 15-JUL-1998 (Rel. 36, Last annotation update)
 DT 15-JUL-1998 (Rel. 36, Last annotation update)
 DE HYPOTHEICAL 60.1 KDA PROTEIN C23C11.06C IN CHROMOSOME 1.
 GN SPAC23C11.06C.
 OS Schizosaccharomyces pombe (Fission yeast).
 CC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
 CC Schizosaccharomycetales; Schizosaccharomycetaceae;
 CC Schizosaccharomycetes.
 OX NCBI_TaxID=4896;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=972;
 RA Brown D., Churcher C.M., Barrell B.G., Rajandream M.A., Wood V.,
 RL Submitted (AUG-1997) to the EMBL/Genbank/DBJ databases.
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration

between the Swiss Institute of Bioinformatics and the EMBL outstation -
the European Bioinformatics Institute. There are no restrictions on its
use by non-profit institutions as long as its content is in no way
modified and this statement is not removed. Usage by and for commercial
entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
or send an email to license@isb-sib.ch).

DR EMBL: 298559; CAB1159.1; -
KW Hypothetical protein; Transmembrane.
FT TRANSMEM 55 75 POTENTIAL.
FT TRANSMEM 82 102 POTENTIAL.
FT TRANSMEM 115 135 POTENTIAL.
FT TRANSMEM 143 163 POTENTIAL.
FT TRANSMEM 201 221 POTENTIAL.
FT TRANSMEM 346 366 POTENTIAL.
SQ SEQUENCE 535 AA; 60124 MW; A6AE149AA2929E2 CRC64;

Query Match 51.2%; Score 44; DB 1; Length 535;
Best Local Similarity 50.0%; Pred. No. 32;
Matches 6; Conservative 1; Mismatches 1; Indels 4; Gaps 1;

OY 4 WPMW----WPMWR 1
DB 183 WSMSPSTWPMRQ 194

RESULT 13
MML4_MYCTU STANDARD; PRT; 967 AA.
ID MML4_MYCTU
AC 053735;
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE PUTATIVE MEMBRANE PROTEIN MML4.
GN MML4 OR RV0450C OR MT0466 OR MT037.14C.
OS Mycobacterium tuberculosis.
OC Bacteria; Firmicutes; Actinobacteriae; Actinobacteridae;
OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1773;
[1]
RN SEQUENCE FROM N.A.
RP STRAIN=H37RV;
RC MEDLINE=98295987; PubMed=9634230;
RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
Gordon S.V., Eigmeier K., Gas S., Barry C.E. III, Tekala F.,
Badoock K., Basham D., Brown D., Chillingworth T., Connor R.,
Davies R., Devlin K., Felwell T., Gentles S., Hamlin N., Holroyd S.,
Hornsbey T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
Rutter S., Seeger K., Skelton S., Squares S., Squares R.,
Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;
RA "Deciphering the biology of Mycobacterium tuberculosis from the
RT complete genome sequence."
RL Nature 393:537-544(1998).
[2]
RN SEQUENCE FROM N.A.
RP STRAIN=CDC 1551 / Oshkosh;
RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
Peterson J., Deboy R., Dodson R., Gwinn M.L., Haft D., Hickey E.,
Kolony J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L.,
Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A.,
Bisai W.;
RA "Whole genome comparison of Mycobacterium tuberculosis clinical and
RT laboratory strains."
Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
-1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
-1- SIMILARITY: BELONGS TO THE MML FAMILY.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
between the Swiss Institute of Bioinformatics and the EMBL outstation,
the European Bioinformatics Institute. There are no restrictions on its
use by non-profit institutions as long as its content is in no way

modified and this statement is not removed. Usage by and for commercial
entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
or send an email to license@isb-sib.ch).

DR EMBL: AL021932; CA17407.1; -
DR EMBL: AE006949; AAK4689.1; -
DR TIGR: MT0466; -
KW Tuberculosis; RV0450C;
DR Hypothetical protein; Transmembrane; Complete proteome.
FT TRANSMEM 26 46 POTENTIAL.
FT TRANSMEM 210 230 POTENTIAL.
FT TRANSMEM 242 262 POTENTIAL.
FT TRANSMEM 303 323 POTENTIAL.
FT TRANSMEM 333 353 POTENTIAL.
FT TRANSMEM 384 404 POTENTIAL.
FT TRANSMEM 769 789 POTENTIAL.
FT TRANSMEM 793 813 POTENTIAL.
FT TRANSMEM 821 841 POTENTIAL.
FT TRANSMEM 875 895 POTENTIAL.
FT TRANSMEM 896 916 POTENTIAL.
SQ SEQUENCE 967 AA; 105234 MW; 6301014031480484 CRC64;

Query Match 51.2%; Score 44; DB 1; Length 967;
Best Local Similarity 75.0%; Pred. No. 55;
Matches 6; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 3 RMPWMPWR 10
DB 930 RMPWMPWR 937

RESULT 14
MML2_MYCTU STANDARD; PRT; 968 AA.
ID MML2_MYCTU
AC 011171;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE PUTATIVE MEMBRANE PROTEIN MML2.
GN MML2 OR RV0507 OR MT0528 OR MT0209.34.
OS Mycobacterium tuberculosis.
OC Bacteria; Firmicutes; Actinobacteriae; Actinobacteridae;
OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1773;
[1]
RN SEQUENCE FROM N.A.
RP STRAIN=H37RV;
RC MEDLINE=98295987; PubMed=9634230;
RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
Gordon S.V., Eigmeier K., Gas S., Barry C.E. III, Tekala F.,
Badoock K., Basham D., Brown D., Chillingworth T., Connor R.,
Davies R., Devlin K., Felwell T., Gentles S., Hamlin N., Holroyd S.,
Hornsbey T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
Rutter S., Seeger K., Skelton S., Squares S., Squares R.,
Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;
RA "Deciphering the biology of Mycobacterium tuberculosis from the
RT complete genome sequence."
RL Nature 393:537-544(1998).
[2]
RN SEQUENCE FROM N.A.
RP STRAIN=CDC 1551 / Oshkosh;
RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
Peterson J., Deboy R., Dodson R., Gwinn M.L., Haft D., Hickey E.,
Kolony J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L.,
Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A.,
Bisai W.;
RA "Whole genome comparison of Mycobacterium tuberculosis clinical and
RT laboratory strains."
Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
-1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
-1- SIMILARITY: BELONGS TO THE MML FAMILY.
CC

CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL: Z77162; CAB00933.1; -
DR EMBL: AE006953; AAK44751.1; -
DR TIGR: MT0528; -
DR Tubercullist: RV0507; -
KW Hypothetical protein: Transmembrane; Complete proteome.
FT TRANSMEM 22 42 POTENTIAL.
FT TRANSMEM 204 224 POTENTIAL.
FT TRANSMEM 245 265 POTENTIAL.
FT TRANSMEM 297 317 POTENTIAL.
FT TRANSMEM 328 348 POTENTIAL.
FT TRANSMEM 378 398 POTENTIAL.
FT TRANSMEM 763 783 POTENTIAL.
FT TRANSMEM 787 807 POTENTIAL.
FT TRANSMEM 815 835 POTENTIAL.
FT TRANSMEM 866 886 POTENTIAL.
FT TRANSMEM 891 911 POTENTIAL.
FT CONFLICT 426 426 R -> H (IN REF. 2).
FT CONFLICT 656 656 E -> A (IN REF. 2).
SQ SEQUENCE 968 AA; 106201 MW; B68AB9B78164EDC0 CRC64;

Query Match 51.2%; Score 44; DB 1; Length 968;
Best Local Similarity 75.0%; Pred. No. 55;
Matches 6; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 3 RWPWPWR 10
|||
DB 924 RWPWPWR 931

RESULT 15
SX13_MOUSE STANDARD; PRT; 984 AA.
ID SX13_MOUSE
AC 004891;
DT 01-JUN-1994 (Rel. 29, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE SOX-13 PROTEIN.
GN SOX13 OR SOX-13.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxId=10090;
RN [1]
RN SEQUENCE FROM N.A., AND FUNCTION (ISOFORM 1).
RC TISSUE=Embryo;
RX MEDLINE=98083175; PubMed=9421502;
RA Roose J., Korver W., Oving E., Wilson A., Wagenaar G., Markman M.,
RA Lamers W., Clevers H.,
RT "High expression of the HMG box factor sox-13 in arterial walls during
RT embryonic development."
RL Nucleic Acids Res. 26:469-476(1998).
RN [2]
RN SEQUENCE FROM N.A., AND FUNCTION (ISOFORM 2).
RC TISSUE=Embryo;
RX MEDLINE=98201614; PubMed=9524265;
RA Kido S., Hirata Y., Ogawa M., Sakai Y., Yoshimura Y., Aiso S.,
RT "Cloning and characterization of mouse msox13 cDNA."
RL Gene 208:201-206(1998).
RN [3]
RN SEQUENCE OF 405-460 FROM N.A.
RX MEDLINE=93181275; PubMed=8441686;
RA Wright E.M., Snopce B., Koopman P.,
RT "Seven new members of the Sox gene family expressed during mouse

RT development."
RL Nucleic Acids Res. 21:744-744(1993).
CC -1- FUNCTION: BINDS TO THE SEQUENCE 5'-AACAT-3'.
CC -1- SUBCELLULAR LOCATION: NUCLEAR (POTENTIAL).
CC -1- ALTERNATIVE PRODUCTS: 2 ISOFORMS; ISOFORM 1 (SHOWN HERE) AND
CC ISOFORM 2; MAY BE PRODUCED BY ALTERNATIVE SPLICING.
CC -1- TISSUE SPECIFICITY: IN THE EMBRYO, HIGH LEVELS OF EXPRESSION ARE
CC FOUND IN THE ARTERIAL WALLS AT 13.5 DAYS POST COITUM (DPC). LOW
CC LEVELS ARE FOUND IN THE INNER EAR AT 13.5 DPC AND IN SOME CELLS IN
CC THE THYMUS AT 16.5 DPC. EXPRESSED IN THE TRACHEAL EPITHELIUM BELOW
CC THE VOCAL CORD AND IN THE HAIR FOLLICLES AT 18 DPC.
CC -1- SIMILARITY: CONTAINS 1 HMG BOX.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL: AJ000740; CA04278.1; -
DR EMBL: AB006329; BAA25786.1; -
DR EMBL: Z18962; CAA79487.1; -
DR PIR: S30241; S30241.
DR HSSP: Q05066; 1HRZ.
DR MGP: MGI:98361; Sox13.
DR InterPro: IPR000910; HMG_12_box.
DR Pfam: PF00505; HMG_box; 2.
DR SMART: SM00398; HMG; 1.
KW DNA-binding; Nuclear protein; Alternative splicing.
FT DOMAIN 159 195 GLN-RICH.
FT DNA_BIND 397 465 HMG_BOX.
FT VARSPLIC 495 519
FT VARSPLIC 603 609 (IN ISOFORM 2).
FT VARSPLIC 610 984 SMWSQT -> ELVVLTQ (IN ISOFORM 2).
FT CONFLICT 35 35 MISSING (IN ISOFORM 2).
FT CONFLICT 41 42 P -> L (IN REF. 2).
FT CONFLICT 195 195 AT -> TN (IN REF. 2).
FT CONFLICT 195 195 AT -> QQ (IN REF. 2).
SQ SEQUENCE 984 AA; 108897 MW; 7F5506EDADEB98C5 CRC64;

Query Match 51.2%; Score 44; DB 1; Length 984;
Best Local Similarity 42.9%; Pred. No. 56;
Matches 6; Conservative 0; Mismatches 0; Indels 8; Gaps 1;

OY 4 WPMW-----PW 9
|||
DB 686 WPMWTKLAEFGSPW 699

Search completed: January 4, 2002, 08:47:48
Job time: 406 sec

THIS PAGE BLANK (USPTO)

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:41:32 ; Search time 27.18 seconds
(without alignments)
33.631 Million cell updates/sec

Title: US-09-444-281-36
Perfect score: 86
Sequence: 1 ILRMPWMPWRRK 12

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 219241 seqs, 76174552 residues

Total number of hits satisfying chosen parameters: 219241

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: PIR:68:*
2: PIR1:*
3: PIR2:*
4: PIR3:*
5: PIR4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	70	81.4	144	1 JCI222	indolicidin precursor
2	53	61.6	1173	1 VG1HMC	E2 glycoprotein pr
3	51	59.3	299	2 T12505	hypothetical prote
4	50	58.1	111	2 T29295	hypothetical prote
5	49	57.0	198	1 J70751	ferredoxin-NADP+
6	48.5	56.4	114	2 T36208	hypothetical prote
7	47	54.7	248	2 S23449	NADH oxidase (H2O2
8	47	54.7	253	2 G70715	hypothetical prote
9	47	54.7	276	2 B83161	probable short-cha
10	47	54.7	715	2 B70741	probable moef prot
11	47	54.7	1411	2 T48529	hypothetical prote
12	46	53.5	728	2 T51071	related to trifa pr
13	45.5	52.9	505	2 A39128	anthranilate synth
14	45	52.3	196	2 S55483	modulator of drug
15	45	52.3	273	2 F82646	monofunctional dio
16	45	52.3	412	2 A83604	probable MFS trans
17	45	52.3	448	2 H72376	hypothetical prote
18	45	52.3	1108	2 A48508	cyclic-nucleotide
19	44	51.2	257	2 S70177	yife protein - Yer
20	44	51.2	361	2 A36669	galactoside 3(4)-L
21	44	51.2	397	2 B70763	probable membrane
22	44	51.2	535	2 T38244	hypothetical prote
23	44	51.2	621	2 S37664	peplomerie polypor
24	44	51.2	630	2 S37663	peplomerie polypor
25	44	51.2	967	2 C70831	probable mmp14 pro
26	44	51.2	968	2 F70746	probable mmp14 pro
27	44	51.2	968	2 T00322	hypothetical prote
28	44	51.2	1134	1 VG1HIB	E2 glycoprotein pr
29	44	51.2	1162	1 VG1HAK	E2 glycoprotein pr

30	44	51.2	1162	2 S07421	E2 glycoprotein pr
31	44	51.2	1162	2 S14939	E2 glycoprotein pr
32	44	51.2	1162	2 S14940	E2 glycoprotein pr
33	43.5	50.6	1529	2 A59189	ATP-binding caset
34	43	50.0	51	2 S23291	light-harvesting p
35	43	50.0	192	2 H86543	hypothetical prote
36	43	50.0	236	2 D72081	conserved hypotet
37	43	50.0	326	2 J00606	arylesterase (EC 3
38	43	50.0	250	2 A83506	probable cobalamin
39	43	50.0	278	2 T46458	hypothetical prote
40	43	50.0	298	2 B72452	hypothetical prote
41	43	50.0	646	2 H82555	c-type cytochrome
42	43	50.0	711	2 C40046	antibiotic transpo
43	43	50.0	738	2 F96701	hypothetical prote
44	43	50.0	958	2 A70634	probable mmp14 pro
45	43	50.0	1112	2 S70522	cyclic nucleotide

ALIGNMENTS

RESULT 1
JCI222
indolicidin precursor - bovine
N:Alternate names: antimicrobial peptide
C:Species: Bos primigenius taurus (cattle)
C>Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 10-Sep-1999
C:Accession: JCI222; A42387; S25664
R:del Sal, G.; Storici, P.; Schneider, C.; Romeo, D.; Zanetti, M.
Biochem. Biophys. Res. Commun. 187, 467-472, 1992
A:Title: cDNA cloning of the neutrophil bactericidal peptide indolicidin.
A:Reference number: JCI222; MUID:92392368
A:Accession: JCI222
A:Molecule type: mRNA
A:Residues: 1-144 <SAL>
A:Cross-references: EMBL: X67340; NID: 9462; PID: CAA47755.1; PID: 9463
A:Experimental source: Bone marrow
R:Seidman, M.E.; Novotny, M.J.; Morris, W.L.; Tang, Y.Q.; Smith, W.; Cullor, J.S.
J. Biol. Chem. 267, 4292-4295, 1992
A:Title: Indolicidin, a novel bactericidal tridecapeptide amide from neutrophils.
A:Reference number: A42387; MUID:92165771
A:Accession: A42387
A:Residues: 131-143 <SEL>
A:Molecule type: protein
A:Experimental source: neutrophils
A>Note: sequence extracted from NCBI backbone (NCBI:83840)
C:Superfamily: catenilin; cystatin homology
C:Keywords: amidated carboxyl end
F:1-29/Domain: signal sequence #status predicted <SIG>
F:32-129/Domain: cystatin homology <CYS>
F:30-130/Domain: propeptide #status predicted <PRO>
F:131-143/Product: indolicidin #status experimental <MAT>
F:143/Modified site: amidated carboxyl end (Arg) (amide in mature form from following

Query Match 81.4%; Score 70; DB 1; Length 144;
Best Local Similarity 88.9%; Pred. No. 0.011;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 3 RMPWMPWRR 11
DB 135 RMPWMPWRR 143

RESULT 2
VG1HMC
E2 glycoprotein precursor - human coronavirus (strain 229E)
N:Alternate names: peplomer glycoprotein; S glycoprotein; spike glycoprotein
C:Species: human coronavirus
A>Note: host Homo sapiens (man)
C>Date: 31-Dec-1991 #sequence_revision 31-Dec-1991 #text_change 16-Jun-2000
C:Accession: A34766; S05460
R:Raabe, T.; Schelle-Prinz, B.; Siddell, S.G.

J. Gen. Virol. 71, 1065-1073, 1990
 A:Title: Nucleotide sequence of the gene encoding the spike glycoprotein of human corona
 A:Reference number: A34766; MUID:90264837
 A:Accession: A34766
 A:Molecule type: mRNA
 A:Residues: 1-1173 <RA>
 A:Cross-references: EMBL:X16816; NID:958926; PTDN:CAA34723.1; PID:958927
 A:Experimental source: strain 229E
 R:Raabe, T.; Sidde11, S.
 Nucleic Acid Res. 17, 6387, 1989
 A:Title: Nucleotide sequence of the human coronavirus HCV 229E mRNA 4 and mRNA 5 unique
 A:Reference number: A34038; MUID:89366667
 A:Accession: S05460
 A:Status: translation not shown
 A:Molecule type: mRNA
 A:Residues: 1159-1173 <RA>
 A:Cross-references: EMBL:X15654; NID:958921; PTDN:CAA33680.1; PID:91334827
 C:Superfamily: coronavirus E2 glycoprotein
 C:Keywords: glycoprotein; transmembrane protein
 F:1-15/Domain: signal sequence #status predicted <SIG>
 F:16-1173/Product: E2 glycoprotein #status predicted <MAT>
 F:1116-1138/Domain: transmembrane #status predicted <TM>
 F:23,62,98,147,171,176,220,243,326,333,440,464,518,538,542,568,581,587,663,671,930,1015,
 E:23,62,98,147,171,176,220,243,326,333,440,464,518,538,542,568,581,587,663,671,930,1015,

Query Match 61.6%; Score 53; DB 1; Length 1173;
 Best Local Similarity 62.5%; Pred. No. 12;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 LRMPWPMW 9
 Db 1112 IKMPWPMW 1119

RESULT 3
 T12505
 hypothetical protein DKFZp434C192.1 - human (fragment)
 C:Species: Homo sapiens (man)
 C:Date: 23-Jul-1999 #sequence_revision 23-Jul-1999 #text_change 23-Jul-1999
 C:Accession: T12505
 R:Ansorge, W.; Wilkner, U.; Mewes, H.W.; Gassenhuber, J.; Wiemann, S.
 submitted to the Protein Sequence Database, June 1999
 A:Reference number: 217527
 A:Accession: T12505
 A:Status: preliminary
 A:Molecule type: mRNA
 A:Residues: 1-299 <ANS>
 A:Cross-references: EMBL:A1096753
 A:Experimental source: adult testis; clone DKFZp434C192
 C:Genetics:
 A:Note: DKFZp434C192.1

Query Match 59.8%; Score 51; DB 2; Length 299;
 Best Local Similarity 85.7%; Pred. No. 6.1;
 Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 5 PMPWPMR 11
 Db 37 PMPWPMR 43

RESULT 4
 T29295
 hypothetical protein C50F7.8 - Caenorhabditis elegans
 C:Species: Caenorhabditis elegans
 C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999
 C:Accession: T29295
 R:Johnson, D.; Steilys, L.
 submitted to the EMBL Data Library, November 1995
 A:Description: The sequence of C. elegans cosmid C50F7.
 A:Reference number: Z20601
 A:Accession: T29295

A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-111 <JOH>
 A:Cross-references: EMBL:U41557; PTDN:AAA83303.1; CESP:C50F7.8
 C:Genetics:
 A:Gene: CESP:C50F7.8

Query Match 58.1%; Score 50; DB 2; Length 111;
 Best Local Similarity 54.5%; Pred. No. 3.2;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 1 LRMPWPMW 11
 Db 12 VMWPMWPGCR 22

RESULT 5
 JF0751
 ferredoxin-NADP+ reductase (EC 1.18.1.2), long form precursor - bovine
 N:Alternate names: adrenodoxin reductase
 C:Species: Bos primigenius taurus (cattle)
 C:Date: 14-Jul-1994 #sequence_revision 18-Oct-1996 #text_change 16-Jun-2000
 C:Accession: JF0751; JF0079; JS0390; S03558; PS0003; A29604; S52100
 R:Takata, Y.; Sagara, Y.; Kono, A.; Sekimizu, K.; Horiiuchi, T.
 Biol. Pharm. Bull. 16, 1200-1206, 1993
 A:Title: Gene structure of bovine adrenodoxin reductase.
 A:Reference number: JF0751; MUID:94177140
 A:Accession: JF0751
 A:Molecule type: DNA
 A:Residues: 1-498 <TA>
 A:Cross-references: GB:D83475; NID:q1199916; PTDN:BA11921.1; PID:94521308
 A:Experimental source: adrenal cortex
 A:Note: the authors translated the codon GNC for residue 205 as Gly
 R:Sagara, Y.; Takata, Y.; Miyata, T.; Hara, T.; Horiiuchi, T.
 J. Biochem. 102, 1333-1336, 1987
 A:Title: Cloning and sequence analysis of adrenodoxin reductase cDNA from bovine adre
 A:Reference number: JF0079; MUID:8818050
 A:Accession: JF0079
 A:Molecule type: mRNA
 A:Residues: 1-204,211-498 <SAG>
 A:Cross-references: GB:D00211; NID:q217433; PTDN:BA00150.1; PID:q217434
 A:Note: the deduced sequence is partially confirmed by amino acid sequencing of 15 is
 R:Sagara, Y.
 submitted to DDBJ, September 1989
 A:Reference number: JS0390
 A:Contents: revision, insertion of residues 205-210
 A:Accession: JS0390
 A:Molecule type: mRNA
 A:Residues: 56-498 <SAG>
 R:Hankoglu, I.; Gutfinger, T.
 Eur. J. Biochem. 180, 479-484, 1989
 A:Title: cDNA sequence of adrenodoxin reductase. Identification of NADP-binding sites
 A:Reference number: S03558; MUID:89170752
 A:Accession: S03558
 A:Molecule type: mRNA
 A:Residues: 135-204,211-498 <HAN>
 A:Cross-references: EMBL:X13736; NID:965; PTDN:CAA32002.1; PID:9833776
 A:Note: 405-Ser was also found
 R:Hamamoto, I.; Kurokohchi, K.; Tanaka, S.; Ichikawa, Y.
 Biochim. Biophys. Acta 953, 207-213, 1988
 A:Title: Adrenoferritin-binding peptide of NADPH-adrenoferritin reductase.
 A:Reference number: PS0003; MUID:88184054
 A:Accession: PS0003
 A:Molecule type: protein
 A:Residues: 33-41, 'S', '43-62:260-283, 'TM', '496-498 <HAM>
 A:Note: a cyanogen bromide peptide binds to adrenoferritin
 R:Nonaka, Y.; Murakami, H.; Yabuchi, Y.; Kuramitsu, S.; Kagamiyama, H.; Yamano, T.;
 Biochem. Biophys. Res. Commun. 145, 1239-1247, 1987
 A:Title: Molecular cloning and sequence analysis of full-length cDNA for adre
 A:Reference number: A29604; MUID:87270696
 A:Accession: A29604
 A:Molecule type: mRNA

A:Residues: 1-76, 'R', 78-80, 'VMALTPRSRL', 95-123, 'RVYRLT', 129-204, 211-273, 'R', 275-322,
A:Cross-references: GB:M17029; NID:g162628; PIDN:AAA30362.1; PID:g162629
A:Experimental source: adrenal cortex
R:Warburton, R.J.; Seybert, D.W.
Biochim. Biophys. Acta 1246, 39-46, 1995
A:Title: Structural and functional characterization of bovine adrenodoxin reductase by 1
A:Reference number: S52100; MUID:95110846
A:Accession: S52100
A:Status: preliminary
A:Molecule type: protein
A:Residues: 'X', 34-41, 'X', 43-48, 'X', 50-51, 304-306, 'X', 308-309, 'X', 311-326 <WAR>
C:Comment: Ferredoxin-NADP+ reductase is localized in the matrix of adrenal cortex mito
ferredoxin-NADP+ reductase, adrenodoxin and two forms of cytochrome P-450.
C:Genetics:
A:Introns: 27/1; 59/3; 91/3; 132/3; 170/3; 204/3; 246/3; 275/1; 341/3; 399/1; 456/1
C:Function:
A:Description: catalyzes the reversible reduction of NADP+ by reduced ferredoxin or red
C:Superfamily: human ferredoxin-NADP+ reductase
C:Keywords: alternative splicing; flavoprotein; mitochondrion; monomer; NADP; oxidoreduc
F:1-32/Domain: transit peptide (mitochondrion) #status predicted <SIG>
F:33-498/Product: ferredoxin-NADP+ reductase, long form #status predicted <MAT>
F:33-204, 211-498/Product: ferredoxin-NADP+ reductase, short form #status experimental <
F:40-70/Region: Delta-alpha-beta FAD nucleotide-binding fold
F:180-190/Region: NADP binding #status predicted
F:281/Binding site: substrate (Lys) #status experimental

Query Match 57.08; Score 49; DB 1; Length 498;
Best Local Similarity 83.38; Pred. No. 18;
Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 4 PWMPPW 9
1 1111
Db 6 WRMPW 11

RESULT 6
136208
hypothetical protein SCE36.09 - Streptomyces coelicolor
C:Species: Streptomyces coelicolor
C:Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 03-Dec-1999
C:Accession: J36208
R:Oliver, K.; Harris, D.; Bentley, S.D.; Parkhill, J.; Barrell, B.G.; Rajandream, M.A.
submitted to the EMBL Data Library, May 1999
A:Reference number: Z21601
A:Accession: J36208
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-114 <OLI>
A:Cross-references: EMBL:AL049763; PIDN:CAB42078.1; GSPDB:GN00070; SCOEDB:SCE36.09
C:Genetics:
A:Gene: SCOEDB:SCE36.09

Query Match 56.48; Score 48.5; DB 2; Length 114;
Best Local Similarity 80.08; Pred. No. 5.1;
Matches 8; Conservative 0; Mismatches 1; Indels 1; Gaps 1;

QY 3 RW-PMWPMR 11
1 111111
Db 103 RWRPMRPMR 112

RESULT 7
S23449
NADH oxidase (H2O2-forming) (EC 1.6.1.1) - Thermus aquaticus
C:Species: Thermus aquaticus
C:Date: 22-Jan-1993 #sequence_revision 22-Jan-1993 #text_change 23-Mar-1993
C:Accession: S23449; S24556
R:Park, H.U.; Kreutzer, R.; Reiser, C.O.A.; Sprinzl, M.
Eur. J. Biochem. 205, 875-879, 1992
A:Title: Molecular cloning and nucleotide sequence of the gene encoding a H(2)O(2)-formi

A:Reference number: S23449; MUID:92249331
A:Accession: S23449
A:Molecule type: DNA
A:Residues: 1-248 <PAR>
A:Cross-references: EMBL:X60110
A:Accession: S24556
A:Molecule type: protein
A:Residues: 1-32 <PAR1>
C:Genetics:
A:Gene: nox
C:Keywords: NAD; oxidoreductase
F:1-248/Product: NADH oxidase (H2O2-forming) #status experimental <MAT>

Query Match 54.78; Score 47; DB 2; Length 248;
Best Local Similarity 100.08; Pred. No. 17;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 5 PWMPPW 9
1 1111
Db 179 PMWPM 183

RESULT 8
G70715
hypothetical protein RV0945 - Mycobacterium tuberculosis (strain H37RV)
C:Species: Mycobacterium tuberculosis
C:Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 20-Jun-2000
C:Accession: G70715
R:Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon
; Connor, R.; Davies, R.; Devlin, K.; Fellwell, T.; Gentles, S.; Hamlin, N.; Holroyd,
Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.
Nature 393, 537-544, 1998
A:Authors: Squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.
A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete geno
A:Reference number: A70500; MUID:98295987
A:Accession: G70715
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-253 <COL>
A:Cross-references: GB:279700; GB:AL123456; NID:g3261628; PIDN:CAB02005.1; PID:g15242
A:Experimental source: strain H37RV
C:Genetics:
A:Gene: RV0945
C:Superfamily: ribitol dehydrogenase: short-chain alcohol dehydrogenase homology
F:8-190/Domain: short-chain alcohol dehydrogenase homology <SADH>

Query Match 54.78; Score 47; DB 2; Length 253;
Best Local Similarity 100.08; Pred. No. 17;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 PWMPPW 9
1 1111
Db 230 PMWPM 234

RESULT 9
B83161
probable short-chain dehydrogenase PA3883 [imported] - Pseudomonas aeruginosa (strain
C:Species: Pseudomonas aeruginosa
C:Date: 15-Sep-2000 #sequence_revision 15-Sep-2000 #text_change 31-Dec-2000
C:Accession: B83161
R:Stover, C.K.; Pham, X.O.; Erwin, A.L.; Mizoguchi, S.D.; Warrenner, P.; Hickey, M.J.;
adman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.; Lardig, K.; L
; Lory, S.; Olson, M.V.
Nature 406, 959-964, 2000
A:Title: Complete genome sequence of Pseudomonas aeruginosa PA01, an opportunistic pa
A:Reference number: A82950; MUID:20437337
A:Accession: B83161
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-276 <STO>

A:Cross-references: GB:AE004805; GB:AE004091; NID:g950055; PIDN:AG07270.1; GSPDB:GN001
 A:Experimental source: strata PA01
 C:Genetics:
 A:Gene: PA3883
 C:Superfamily: ribitol dehydrogenase; short-chain alcohol dehydrogenase homology

Query Match 54.7%; Score 47; DB 2; Length 276;
 Best Local Similarity 60.0%; Pred. No. 18;
 Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

OY 3 RMPWMPRRK 12
 Db 197 RSPWMPLRQ 206

RESULT 10
 B70741
 probable moey protein - Mycobacterium tuberculosis (strain H37RV)

C:Species: Mycobacterium tuberculosis
 C:Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 22-Oct-1999
 C:Accession: B70741
 R:Coile, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.; Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd, S.; Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.
 A:Authors: Squires, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.
 A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete genome
 A:Reference number: A70500; MUID:98295987
 A:Accession: B70741
 A:Status: preliminary; nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-715 <COL>
 A:Cross-references: GB:275555; GB:AL123456; NID:g3261608; PIDN:CA99988.1; PID:e250356;
 A:Experimental source: strain H37RV
 C:Genetics:
 A:Gene: moey

Query Match 54.7%; Score 47; DB 2; Length 715;
 Best Local Similarity 66.7%; Pred. No. 46;
 Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 3 RMPWMPRRK 11
 Db 65 RWAYTPMRK 73

RESULT 11
 T48529
 hypothetical protein T22P22.90 - Arabidopsis thaliana
 C:Species: Arabidopsis thaliana (mouse-ear cress)
 C:Date: 20-Apr-2000 #sequence_revision 20-Apr-2000 #text_change 20-Apr-2000
 C:Accession: T48529
 R:Bevan, M.; Hilbert, H.; Braun, M.; Holzer, E.; Brandt, A.; Duesterhoft, A.; Bancroft, submitted to the protein Sequence Database, April 2000
 A:Reference number: Z24450
 A:Accession: T48529
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-1411 <BEV>
 A:Cross-references: EMBL:AL163814
 A:Experimental source: cultivar Columbia; BAC clone T22P22
 C:Genetics:
 A:Map position: 5
 A:Introns: 281/2; 320/1; 389/3; 429/3; 473/3; 515/3; 534/2; 567/3; 602/1; 669/1; 776/2;
 A:Note: T22P22.90

Query Match 54.7%; Score 47; DB 2; Length 1411;
 Best Local Similarity 63.6%; Pred. No. 87;
 Matches 7; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

OY 2 LRMPWMPRRK 12
 Db 1013 LAMSWQWRRK 1023

RESULT 12
 T51071
 related to trfA protein [imported] - Neurospora crassa

N:Alternate names: protein B2A19.50
 C:Species: Neurospora crassa
 C:Date: 21-Jul-2000 #sequence_revision 21-Jul-2000 #text_change 21-Jul-2000
 C:Accession: T51071
 R:Schulte, U.; Aign, V.; Hohnsels, J.; Brandt, P.; Fartmann, B.; Holland, R.; Nyakatu submitted to the Protein Sequence Database, July 2000
 A:Reference number: Z25286
 A:Accession: T51071
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-728 <SCH>
 A:Cross-references: EMBL:AL390092; GSPDB:GN00116; NCSP:B2A19.50
 A:Experimental source: BAC clone B2A19; strain OR74A
 C:Genetics:
 A:Gene: NCSP:B2A19.50
 A:Map position: 6
 A:Introns: 26/1; 119/2

Query Match 53.5%; Score 46; DB 2; Length 728;
 Best Local Similarity 58.3%; Pred. No. 62;
 Matches 7; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

OY 1 ILRWMPWRRK 12
 Db 11 ILGWPLWMSRR 22

RESULT 13
 A39128
 anthranilate synthase (EC 4.1.3.27) component I [validated] - Pseudomonas syringae pv
 N:Alternate names: anthranilate synthase alpha chain
 C:Species: Pseudomonas syringae pv. savastanoi
 C:Date: 27-Nov-1991 #sequence_revision 27-Nov-1991 #text_change 17-Mar-2000
 C:Accession: A39128
 R:da Costa, E.; Silva, O.; Kosuge, T.
 J. Bacteriol. 173, 463-471, 1991
 A:Title: Molecular characterization and expression analysis of the anthranilate synth
 A:Reference number: A39128; MUID:91100331
 A:Accession: A39128
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-505 <DAC>
 A:Cross-references: GB:M55911
 C:Genetics:
 A:Gene: type
 C:Complex: heterotrimer; two component I chains, two component II chains
 C:Function: <ANT>
 A:Description: EC 4.1.3.27 [validated; MUID:901303251]
 A:Pathway: tryptophan biosynthesis
 A:Note: first step
 C:Function: <COM1>
 A:Description: EC 4.1.3.27 [validated; MUID:91100331]
 A:Note: expression of trpE seems to be independent of the concentration of tryptophan
 C:Superfamily: anthranilate synthase component I
 C:Keywords: carbon-carbon lyase; oxo-acid-lyase; tryptophan biosynthesis

Query Match 52.9%; Score 45.5; DB 2; Length 505;
 Best Local Similarity 28.0%; Pred. No. 51;
 Matches 7; Conservative 2; Mismatches 3; Indels 13; Gaps 1;

OY 1 ILRW-----PMPWRRK 12
 Db 467 VLEWETLNKRRAMVGSAMPWRR 491

THIS PAGE BLANK (USPTO)

APPLICANT: Kim, Jeong Hyun
APPLICANT: Hong, Seung-Suh
APPLICANT: Lee, Hyun-Soo
APPLICANT: Samsung Genex Corporation
TITLE OF INVENTION: METHOD FOR MASS PRODUCTION OF
FILE REFERENCE: 6181/0F135
CURRENT APPLICATION NUMBER: US/09/230,180
CURRENT FILING DATE: 1999-03-10
PRIOR APPLICATION NUMBER: PCT/KR98/00132
PRIOR FILING DATE: 1998-05-28
PRIOR APPLICATION NUMBER: KR 13372/1998
PRIOR FILING DATE: 1998-04-09
PRIOR APPLICATION NUMBER: KR 21312/1997
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 36
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 29
LENGTH: 39
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: DNA sequence deduced from Indolicidin peptide
OTHER INFORMATION: sequence based on codon usage of E. coli
US-09-230-180-29

alignment_scores:
Quality: 70.00 Length: 9
Ratio: 7.778 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:

US-09-444-281-36 x US-09-230-180-29 ..

Align seg 1/1 to: US-09-230-180-29 from: 1 to: 39

3 ArgTrpProtTrpProtTrpArg 11
:|||||
13 AATGGCGGTGGTGGCGGTGGT 39

seq_name: /cgn2_6/ptodata/2/lna/6A_COMB.seq:US-09-259-741-5

seq_documentation_block:

Sequence 5, Application US/09259741
Patent No. 6033895

GENERAL INFORMATION:

APPLICANT: GARGER, STEPHEN

APPLICANT: HOLTZ, R. BARRY

APPLICANT: MCCULLOCH, MICHAEL

APPLICANT: TURPEN, THOMAS

TITLE OF INVENTION: A PROCESS FOR ISOLATING AND

TITLE OF INVENTION: PURIFYING VIRUSES SOLUBLE PROTEINS AND PEPTIDES FROM PLANT

NUMBER OF SEQUENCES: 5

CORRESPONDENCE ADDRESS:

ADDRESSEE: Howrey & Simon

STREET: 1299 Pennsylvania Avenue N.W.

CITY: Washington

STATE: DC

COUNTRY: USA

ZIP: 20004

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

COMPUTER: IBM Compatible

OPERATING SYSTEM: DOS

SOFTWARE: FastSeq for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/259,741

FILING DATE: February 25, 1999

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 09/037,751
FILING DATE: March 10, 1998
ATTORNEY/AGENT INFORMATION:
NAME: Halluin, Albert P
REGISTRATION NUMBER: 25,277
REFERENCE/DOCKET NUMBER: 00801, 0140, US01
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-463-8100
TELEFAX: 650-463-8400
TELEX:
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 6446 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: unknown
MOLECULE TYPE: Genomic RNA
US-09-259-741-5

alignment_scores:
Quality: 70.00 Length: 9
Ratio: 7.778 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:

US-09-444-281-36 x US-09-259-741-5 ..

Align seg 1/1 to: US-09-259-741-5 from: 1 to: 6446

3 ArgTrpProtTrpProtTrpArg 11
:|||||
6213 AAGUGCCUUGGUGGCCAUGGCGCGA 6239

seq_name: /cgn2_6/ptodata/2/lna/6A_COMB.seq:US-09-037-751-5

seq_documentation_block:

Sequence 5, Application US/09037751
Patent No. 6037456

GENERAL INFORMATION:

APPLICANT: GARGER, STEPHEN

APPLICANT: HOLTZ, R. BARRY

APPLICANT: MCCULLOCH, MICHAEL

APPLICANT: TURPEN, THOMAS

TITLE OF INVENTION: A PROCESS FOR ISOLATING AND

TITLE OF INVENTION: PURIFYING VIRUSES SOLUBLE PROTEINS AND PEPTIDES

NUMBER OF SEQUENCES: 5

CORRESPONDENCE ADDRESS:

ADDRESSEE: Howrey & Simon

STREET: 1299 Pennsylvania Avenue N.W.

CITY: Washington

STATE: DC

COUNTRY: USA

ZIP: 20004

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

COMPUTER: IBM Compatible

OPERATING SYSTEM: DOS

SOFTWARE: FastSeq for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/037,751

FILING DATE: 10-MAR-1998

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER:

FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Halluin, Albert P

REGISTRATION NUMBER: 25,277

REFERENCE/DOCKET NUMBER: 00801, 0140, 999

TELECOMMUNICATION INFORMATION:

TELEPHONE: 650-463-8109

```

1 TELEFAX: 650-463-8400
2
3 TELE:
4
5 INFORMATION FOR SEQ ID NO: 5:
6
7 SEQUENCE CHARACTERISTICS:
8
9 LENGTH: 6446 base pairs
10
11 TYPE: nucleic acid
12
13 STRANDEDNESS: single
14
15 TOPOLOGY: unknown
16
17 MOLECULE TYPE: Genomic RNA
18
19 US-09-037-751-5
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65
66
67
68
69
70
71
72
73
74
75
76
77
78
79
80
81
82
83
84
85
86
87
88
89
90
91
92
93
94
95
96
97
98
99
100
101
102
103
104
105
106
107
108
109
110
111
112
113
114
115
116
117
118
119
120
121
122
123
124
125
126
127
128
129
130
131
132
133
134
135
136
137
138
139
140
141
142
143
144
145
146
147
148
149
150
151
152
153
154
155
156
157
158
159
160
161
162
163
164
165
166
167
168
169
170
171
172
173
174
175
176
177
178
179
180
181
182
183
184
185
186
187
188
189
190
191
192
193
194
195
196
197
198
199
200
201
202
203
204
205
206
207
208
209
210
211
212
213
214
215
216
217
218
219
220
221
222
223
224
225
226
227
228
229
230
231
232
233
234
235
236
237
238
239
240
241
242
243
244
245
246
247
248
249
250
251
252
253
254
255
256
257
258
259
260
261
262
263
264
265
266
267
268
269
270
271
272
273
274
275
276
277
278
279
280
281
282
283
284
285
286
287
288
289
290
291
292
293
294
295
296
297
298
299
300
301
302
303
304
305
306
307
308
309
310
311
312
313
314
315
316
317
318
319
320
321
322
323
324
325
326
327
328
329
330
331
332
333
334
335
336
337
338
339
340
341
342
343
344
345
346
347
348
349
350
351
352
353
354
355
356
357
358
359
360
361
362
363
364
365
366
367
368
369
370
371
372
373
374
375
376
377
378
379
380
381
382
383
384
385
386
387
388
389
390
391
392
393
394
395
396
397
398
399
400
401
402
403
404
405
406
407
408
409
410
411
412
413
414
415
416
417
418
419
420
421
422
423
424
425
426
427
428
429
430
431
432
433
434
435
436
437
438
439
440
441
442
443
444
445
446
447
448
449
450
451
452
453
454
455
456
457
458
459
460
461
462
463
464
465
466
467
468
469
470
471
472
473
474
475
476
477
478
479
480
481
482
483
484
485
486
487
488
489
490
491
492
493
494
495
496
497
498
499
500
501
502
503
504
505
506
507
508
509
510
511
512
513
514
515
516
517
518
519
520
521
522
523
524
525
526
527
528
529
530
531
532
533
534
535
536
537
538
539
540
541
542
543
544
545
546
547
548
549
550
551
552
553
554
555
556
557
558
559
560
561
562
563
564
565
566
567
568
569
570
571
572
573
574
575
576
577
578
579
580
581
582
583
584
585
586
587
588
589
590
591
592
593
594
595
596
597
598
599
600
601
602
603
604
605
606
607
608
609
610
611
612
613
614
615
616
617
618
619
620
621
622
623
624
625
626
627
628
629
630
631
632
633
634
635
636
637
638
639
640
641
642
643
644
645
646
647
648
649
650
651
652
653
654
655
656
657
658
659
660
661
662
663
664
665
666
667
668
669
670
671
672
673
674
675
676
677
678
679
680
681
682
683
684
685
686
687
688
689
690
691
692
693
694
695
696
697
698
699
700
701
702
703
704
705
706
707
708
709
710
711
712
713
714
715
716
717
718
719
720
721
722
723
724
725
726
727
728
729
730
731
732
733
734
735
736
737
738
739
740
741
742
743
744
745
746
747
748
749
750
751
752
753
754
755
756
757
758
759
760
761
762
763
764
765
766
767
768
769
770
771
772
773
774
775
776
777
778
779
780
781
782
783
784
785
786
787
788
789
790
791
792
793
794
795
796
797
798
799
800
801
802
803
804
805
806
807
808
809
810
811
812
813
814
815
816
817
818
819
820
821
822
823
824
825
826
827
828
829
830
831
832
833
834
835
836
837
838
839
840
841
842
843
844
845
846
847
848
849
850
851
852
853
854
855
856
857
858
859
860
861
862
863
864
865
866
867
868
869
870
871
872
873
874
875
876
877
878
879
880
881
882
883
884
885
886
887
888
889
890
891
892
893
894
895
896
897
898
899
900
901
902
903
904
905
906
907
908
909
910
911
912
913
914
915
916
917
918
919
920
921
922
923
924
925
926
927
928
929
930
931
932
933
934
935
936
937
938
939
940
941
942
943
944
945
946
947
948
949
950
951
952
953
954
955
956
957
958
959
960
961
962
963
964
965
966
967
968
969
970
971
972
973
974
975
976
977
978
979
980
981
982
983
984
985
986
987
988
989
990
991
992
993
994
995
996
997
998
999
1000
1001
1002
1003
1004
1005
1006
1007
1008
1009
1010
1011
1012
1013
1014
1015
1016
1017
1018
1019
1020
1021
1022
1023
1024

```

```

; MOLECULE TYPE: Genomic RNA
; SEQUENCE DESCRIPTION: SEQ ID NO: 5 :
US-09-466-422-5

alignment_scores:
      Quality: 70.00          Length: 9
      Ratio: 7.778           Gaps: 0
Percent Similarity: 100.000    Percent Identity: 88.889

alignment_block:
US-09-444-281-36 x US-09-466-422-5 ..

Align seg 1/1   to: US-09-466-422-5   from: 1   to: 6446

      3 ARGTPRPTRTPRTPRTPARGATG 11
      ::::::::::::::::::::
76213 AAGUGCCUUGGCGGCCAUGC GCCCA 62339

seq_name: /cgn2_6/p/tdat01/2/lna/6A-COMB.seq:US-08-793-035-6

seq_documentation_block:
; Sequence 6, Application US/08793035
; Patent No. 6011201
; GENERAL INFORMATION:
; APPLICANT: Slabas, Antoni R.
; APPLICANT: White, Andrew
; APPLICANT: Chase, Dianne
; APPLICANT: Elborough, Keiran
; APPLICANT: Fenem, Phillip A.
; TITLE OF INVENTION: B-ketacyl ACP Reductase Genes From
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Arnold White & Durkee
; STREET: P O. Box 4433
; CITY: Houston
; STATE: TX
; COUNTRY: US
; ZIP: 77210-4433
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0., Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/793,035
; FILING DATE: 28-JUL-1997
; CLASSIFICATION: 800
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9416422.2
; FILING DATE: 20-JUL-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB95/01678
; FILING DATE: 17-JUL-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Kammerer, Patricia A.
; REGISTRATION NUMBER: 29,775
; REFERENCE/DOCKET NUMBER: MOST:132
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 713.787.1400
; TELEFAX: 713.787.1440
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 758 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-793-035-6

alignment_scores:
Quality: 61.00          Length: 9
```

Ratio: 7.625 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 77.778

alignment_block:

US-09-444-281-36 x US-08-793-035-6/rev ..

Align seg 1/1 to reverse of: US-08-793-035-6 from: 1 to: 758

1 l l e u a r g t r p r o t r p r o t r p 9
:
451 CTCCTGCATGAGTGTGAGCTGG 425

seq_name: /cgn2_6/ptodata/2/lna/6B_COMB.seq:US-09-020-956-82

seq_documentation_block:

; Sequence 82, Application US/09020956;
; Patent No. 6261562

GENERAL INFORMATION:

; APPLICANT: Xu, Jiangchun

; APPLICANT: Dillon, David C.

; TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY OF PROSTATE CANCER AND METHODS FO

; NUMBER OF SEQUENCES: 178

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: SEED and BERRY LLP

; STREET: 6300 Columbia Center, 701 Fifth Avenue

; CITY: Seattle

; STATE: WA

; COUNTRY: USA

; ZIP: 98104

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patent In Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; FILING DATE: 09-FEB-1998

; CLASSIFICATION:

; ATTORNEY/AGENT INFORMATION:

; NAME: Maki, David J.

; REGISTRATION NUMBER: 31,392

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (206) 622-4900

; TELEFAX: (206) 682-6031

; INFORMATION FOR SEQ ID NO: 82:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 383 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; MOLECULE TYPE: cDNA

; ORIGINAL SOURCE:

; ORGANISM: Homo sapiens

alignment_scores:

Quality: 59.00 Length: 8

Ratio: 8.429 Gaps: 0

Percent Similarity: 87.500 Percent Identity: 87.500

alignment_block:

US-09-444-281-36 x US-09-020-956-82 ..

Align seg 1/1 to: US-09-020-956-82 from: 1 to: 383

2 l e u a r g t r p r o t r p r o t r p 9
| | | | | | | | | | | | | | | | | | | | | |
155 CTCGCTGCCTGTGAGCTGG 178

seq_name: /cgn2_6/ptodata/2/lna/6B_COMB.seq:US-09-030-607-82

seq_documentation_block:
; Sequence 82, Application US/09030607
; Patent No. 6262245

GENERAL INFORMATION:

; APPLICANT: Xu, Jiangchun

; APPLICANT: Dillon, David C.

; TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY OF PROSTATE CANCER AND METHODS

; NUMBER OF SEQUENCES: 224

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: SEED and BERRY LLP

; STREET: 6300 Columbia Center, 701 Fifth Avenue

; CITY: Seattle

; STATE: WA

; COUNTRY: USA

; ZIP: 98104

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patent In Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/09/030,607

; FILING DATE: 25-FEB-1998

; CLASSIFICATION:

; ATTORNEY/AGENT INFORMATION:

; NAME: Maki, David J.

; REGISTRATION NUMBER: 31,392

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (206) 622-4900

; TELEFAX: (206) 682-6031

; INFORMATION FOR SEQ ID NO: 82:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 383 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; MOLECULE TYPE: cDNA

; ORIGINAL SOURCE:

; ORGANISM: Homo sapiens

alignment_scores:

Quality: 59.00 Length: 8

Ratio: 8.429 Gaps: 0

Percent Similarity: 87.500 Percent Identity: 87.500

alignment_block:

US-09-444-281-36 x US-09-030-607-82 ..

Align seg 1/1 to: US-09-030-607-82 from: 1 to: 383

2 l e u a r g t r p r o t r p r o t r p 9
| | | | | | | | | | | | | | | | | | | | | |
155 CTCGCTGCCTGTGAGCTGG 178

seq_name: /cgn2_6/ptodata/2/lna/6B_COMB.seq:US-09-030-607-183

seq_documentation_block:

; Sequence 183, Application US/09030607
; Patent No. 6262245

GENERAL INFORMATION:

; APPLICANT: Xu, Jiangchun

; APPLICANT: Dillon, David C.

; TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY OF PROSTATE CANCER AND METHODS

; NUMBER OF SEQUENCES: 224

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: SEED and BERRY LLP

; STREET: 6300 Columbia Center, 701 Fifth Avenue

; CITY: Seattle

; STATE: WA

; COUNTRY: USA


```

: ZIP: 98104
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/09/030.607
: FILING DATE: 25-FEB-1998
: CLASSIFICATION:
: ATTORNEY/AGENT INFORMATION:
: NAME: Makl, David J.
: REGISTRATION NUMBER: 31,392
: REFERENCE/DOCKET NUMBER: 210121.427C3
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (206) 622-4900
: TELEFAX: (206) 682-6031
: INFORMATION FOR SEQ ID NO: 183:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 384 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
:
: US-09-030-607-183

alignment_scores:
: Quality: 59.00 Length: 8
: Ratio: 8.429 Gaps: 0
: Percent Similarity: 87.500 Percent Identity: 87.500

alignment_block:
: US-09-444-281-36 x US-09-030-607-183 ..
:
: Align seg 1/1 to: US-09-030-607-183 from: 1 to: 384
:
: 2 LeuArgTrpProTrpPrpTrp 9
: |||||
: 156 CTTCGCTGCGCTGTGAGAGCTGG 179

seq_name: /cgn2_6/ptodata/2/ina/6B_COMB.seq:US-09-020-956-73

seq_documentation_block:
: Sequence 73, Application US/09020956
: Patent No. 6261562
: GENERAL INFORMATION:
: APPLICANT: Xu, Jiangchun
: APPLICANT: Dillon, Davin C.
: TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY OF PROSTATE CANCER AND METHODS FO
: NUMBER OF SEQUENCES: 178
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: SEED and BERRY LLP
: STREET: 6300 Columbia Center, 701 Fifth Avenue
: CITY: Seattle
: STATE: WA
: COUNTRY: USA
: ZIP: 98104
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/09/020.956
: FILING DATE: 09-FEB-1998
: CLASSIFICATION:
: ATTORNEY/AGENT INFORMATION:
: NAME: Makl, David J.
: REGISTRATION NUMBER: 31,392
: REFERENCE/DOCKET NUMBER: 210121.427C2
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (206) 622-4900
: TELEFAX: (206) 682-6031
```

```

: INFORMATION FOR SEQ ID NO: 73:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 499 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: cDNA
: ORIGINAL SOURCE:
: ORGANISM: Homo sapiens
:
: US-09-020-956-73

alignment_scores:
: Quality: 59.00 Length: 8
: Ratio: 8.429 Gaps: 0
: Percent Similarity: 87.500 Percent Identity: 87.500

alignment_block:
: US-09-444-281-36 x US-09-020-956-73 ..
:
: Align seg 1/1 to: US-09-020-956-73 from: 1 to: 499
:
: 2 LeuArgTrpProTrpPrpTrp 9
: |||||
: 115 CTTCGCTGCGCTGTGAGAGCTGG 138

seq_name: /cgn2_6/ptodata/2/ina/6B_COMB.seq:US-09-030-607-73

seq_documentation_block:
: Sequence 73, Application US/09030607
: Patent No. 6262245
: GENERAL INFORMATION:
: APPLICANT: Xu, Jiangchun
: APPLICANT: Dillon, Davin C.
: TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY OF PROSTATE CANCER AND METHODS
: NUMBER OF SEQUENCES: 224
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: SEED and BERRY LLP
: STREET: 6300 Columbia Center, 701 Fifth Avenue
: CITY: Seattle
: STATE: WA
: COUNTRY: USA
: ZIP: 98104
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/09/030.607
: FILING DATE: 25-FEB-1998
: CLASSIFICATION:
: ATTORNEY/AGENT INFORMATION:
: NAME: Makl, David J.
: REGISTRATION NUMBER: 31,392
: REFERENCE/DOCKET NUMBER: 210121.427C3
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (206) 622-4900
: TELEFAX: (206) 682-6031
: INFORMATION FOR SEQ ID NO: 73:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 499 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: cDNA
: ORIGINAL SOURCE:
: ORGANISM: Homo sapiens
:
: US-09-030-607-73

alignment_scores:
: Quality: 59.00 Length: 8
```

Ratio: 8.429 Gaps: 0
Percent Similarity: 87.500 Percent Identity: 87.500

alignment_block:
US-09-444-281-36 x US-09-030-607-73

Align seg 1/1 to: US-09-030-607-73 from: 1 to: 499

2 LeuArgTrpProTrpTrpProTrp 9
|||||
115 CTGCGTGGCCCTGTGTGAGCTGG 138

seq_name: /cgn2_6/ptodata/2/lna/6B_COMB.seq:US-09-088-651-3

seq_documentation_block:

Sequence 3, Application US/09088651
Patent No. 6165771

GENERAL INFORMATION:

APPLICANT: BURGESS, NICOLA A.

APPLICANT: CLINKENBEARD, HELEN E.

APPLICANT: SOUTHAN, CHRISTOPHER D.

TITLE OF INVENTION: NOVEL COMPOUNDS

NUMBER OF SEQUENCES: 6

CORRESPONDENCE ADDRESS:

ADDRESSEE: RATNER & PRESTIA

STREET: P.O. BOX 980

CITY: VALLEY FORGE

STATE: PA

COUNTRY: USA

ZIP: 19482

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

COMPUTER: IBM Compatible

OPERATING SYSTEM: DOS

SOFTWARE: FASTSEQ for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/088,651

FILING DATE: JUNE 1, 1998

CLASSIFICATION:

Prior Application DATA:

APPLICATION NUMBER: GB9712088.5

FILING DATE: 10-JUNE-1997

APPLICATION NUMBER: EP 97308295.1

FILING DATE: 17-OCT-1997

APPLICATION NUMBER: GB 9803650.2

FILING DATE: 20-FEB-1998

ATTORNEY/AGENT INFORMATION:

NAME: PRESTIA, PAUL F.

REGISTRATION NUMBER: 23,031

REFERENCE/DOCKET NUMBER: GH30358

TELECOMMUNICATION INFORMATION:

TELEPHONE: 610-407-0700

TELEFAX: 610-407-0701

TELEX: 846169

INFORMATION FOR SEQ. ID NO. 3:

SEQUENCE CHARACTERISTICS:

LENGTH: 683 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: cDNA

US-09-088-651-3

alignment_scores:

Quality: 59.00 Length: 11

Ratio: 6.556 Gaps: 0

Percent Similarity: 81.818 Percent Identity: 63.636

alignment_block:

US-09-444-281-36 x US-09-088-651-3

Align seg 1/1 to: US-09-088-651-3 from: 1 to: 683

2 LeuArgTrpProTrpTrpProTrpArgLys 12
||| |||||
139 CTGACGTGGCCCTGTGTGAGCTGGAGGCACAG 171

seq_name: /cgn2_6/ptodata/2/lna/6B_COMB.seq:US-09-088-651-1

seq_documentation_block:

Sequence 1, Application US/09088651

Patent No. 6165771

GENERAL INFORMATION:

APPLICANT: BURGESS, NICOLA A.

APPLICANT: CLINKENBEARD, HELEN E.

APPLICANT: SOUTHAN, CHRISTOPHER D.

TITLE OF INVENTION: NOVEL COMPOUNDS

NUMBER OF SEQUENCES: 6

CORRESPONDENCE ADDRESS:

ADDRESSEE: RATNER & PRESTIA

STREET: P.O. BOX 980

CITY: VALLEY FORGE

STATE: PA

COUNTRY: USA

ZIP: 19482

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

COMPUTER: IBM Compatible

OPERATING SYSTEM: DOS

SOFTWARE: FASTSEQ for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/088,651

FILING DATE: JUNE 1, 1998

CLASSIFICATION:

Prior Application DATA:

APPLICATION NUMBER: GB9712088.5

FILING DATE: 10-JUNE-1997

APPLICATION NUMBER: EP 97308295.1

FILING DATE: 17-OCT-1997

APPLICATION NUMBER: GB 9803650.2

FILING DATE: 20-FEB-1998

ATTORNEY/AGENT INFORMATION:

NAME: PRESTIA, PAUL F.

REGISTRATION NUMBER: 23,031

REFERENCE/DOCKET NUMBER: GH30358

TELECOMMUNICATION INFORMATION:

TELEPHONE: 610-407-0700

TELEFAX: 610-407-0701

TELEX: 846169

INFORMATION FOR SEQ. ID NO. 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 1109 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: cDNA

US-09-088-651-1

alignment_scores:

Quality: 59.00 Length: 11

Ratio: 6.556 Gaps: 0

Percent Similarity: 81.818 Percent Identity: 63.636

alignment_block:

US-09-444-281-36 x US-09-088-651-1

Align seg 1/1 to: US-09-088-651-1 from: 1 to: 1109

2 LeuArgTrpProTrpTrpProTrpArgLys 12
||| |||||
564 CTGACGTGGCCCTGTGTGAGCTGGAGGCACAG 596

seq_name: /cgn2_6/ptodata/2/lna/6B_COMB.seq:US-09-088-651-6

```

? APPLICANT : SAMPATH, Kuber T.
? TITLE OF INVENTION : Morphogenic Protein-Specific Cell
? TITLE OF INVENTION : Surface Receptors and Uses Therefor
? NUMBER OF SEQUENCES : 18
? CORRESPONDENCE ADDRESS :
? ADDRESSEE : Testa, Hurwitz & Thibault
? STREET : 125 High St.
? CITY : Boston
? STATE : MA
? COUNTRY : USA
? ZIP : 02110
? COMPUTER READABLE FORM:
? MEDIUM TYPE : Floppy disk
? COMPUTER : IBM PC compatible
? OPERATING SYSTEM : PC-DOS/MS-DOS
? SOFTWARE : Patent Release #1.0, Version #1.30
? CURRENT APPLICATION DATA:
? APPLICATION NUMBER: US/08/481,337A
? FILING DATE : 02-JUN-1995
? CLASSIFICATION : 435
? ATTORNEY/AGENT INFORMATION:
? NAME : MEYERS, Thomas C.
? REGISTRATION NUMBER : 36,989
? REFERENCE/DOCKET NUMBER: CRP-097CP2
? TELECOMMUNICATION INFORMATION:
? TELEPHONE : (617) 248-7000
? TELEFAX : (617) 248-7100
? INFORMATION FOR SEQ ID NO: 1 :
? SEQUENCE CHARACTERISTICS:
? LENGTH: 1509 base pairs
? TYPE: nucleic acid
? STRANDEDNESS: single
? TOPOLOGY: linear
? MOLECULE TYPE: cDNA
? FEATURE:
? NAME/KEY : CDS
? LOCATION : 1..1509
? OTHER INFORMATION : /product= "Human ALK1"
US-08-481-337A-1

alignment_scores:
    Quality: 58.00      Length: 6
    Ratio: 9.667       Gaps: 0
Percent Similarity: 100.000   Percent Identity: 100.000

alignment_block:
US-09-444-281-36 x US-08-481-337A-1 ..

Align seg 1/1 to: US-08-481-337A-1 from: 1 to: 1509

      4 TrpProTrpTrpProTrp 9
        |||||||||
389 TGCCCTGCTGGCCTCG 406

```

THIS PAGE BLANK (USPTO)


```

ORGANISM      Nicotiana benthamiana
               Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
               Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
               Asteridae; euasterids I; Solanales; Solanaceae; Nicotiana.
REFERENCE     1 (bases 1 to 6446)
AUTHORS       Garger,S.J., Holtz,B.R., Mcclulloch,M.J. and Tuppen,T.H.
TITLE         A process for isolating and purifying viruses, soluble proteins and
               peptides from plant sources
               Patent: WO 011969-A 5 22-MAR-2001;
               Large Scale Biology Corporation (US)
FEATURES      Location/Qualifiers
               1..6446
               /organism="Nicotiana benthamiana"
               /db_xref="taxon:4100"
BASE COUNT    1873 a 1234 c 1563 g 1776 t
ORIGIN
alignment_scores:
               Quality: 70.00      Length: 9
               Ratio: 7.778        Gaps: 0
               Percent Similarity: 100.000      Percent Identity: 88.889
alignment_block:
US-09-444-281-36 x AX098418      ..
Align seg 1/1 to: AX098418 from: 1 to: 6446
3 ArgTrpProTrrPrProTrrPARGATG 11
:::|||||
6213 AAGTGGCTTGGTGGCCATGGCCGCA 6239
seq_name: gb_to:MMU08210
seq_documentation_block:
LOCUS      MMU08210      2651 bp      mRNA      ROD      31-OCT-1995
DEFINITION Mus musculus tropoelastin mRNA, complete cds.
ACCESSION  U08210
VERSION     U08210.1 GI:473273
KEYWORDS   .
SOURCE     house mouse..
            Mus musculus
ORGANISM   Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE  1 (bases 1 to 2651)
AUTHORS   Wjdane,K.S., Sechler,J.L., Boyd,C.D. and Passmore,H.C.
TITLE     Use of an inltron polymorphism to localize the tropoelastin gene to
            mouse chromosome 5 in a region of linkage conservation with human
            chromosome 7
            Genomics 23 (1), 125-131 (1994)
JOURNAL    MEDLINE
MEDLINE    95130069
REFERENCE  2 (bases 1 to 2651)
AUTHORS    Boyd,C.D.
TITLE      Direct Submission
JOURNAL    Submitted (30-MAR-1994) Charles D. Boyd, Department of Surgery,
            UMDNJ - Robert Wood Johnson Medical School, 51 French St., New
            Brunswick, NJ 08903, USA
            Location/Qualifiers
            1..2651
            /organism="Mus musculus"
            /strain="BALB/c"
            /db_xref="taxon:10090"
            /chromosome="5"
            /tissue_type="lung"
            /dev_stage="adult"
            1..2583
            /codon_start=1
            /product="tropoelastin"
            /protein_id="AA80155.1"
            /db_xref="GI:473274"
            /translation="MAGITAVVDPQPGVLLILNLTLPADPGVPAVPGGLPGGVG
            GVVYPACIGIGIGGGGALGPCKPPKPGAGLICTFAGPGGLGAGPGAGLGAFFPGV
            TFGAGGLVYGGAAGAAAYKAAKKAGAGLGVGVGVGVGVGVGPGVGVGVGVG"
CDS

```

BASE COUNT	410 a	623 c	1029 g	589 t
ORIGIN	RRR"			
alignment_scores:				
Quality:	69.00	Length:	10	
Ratio:	7.667	Gaps:	0	
Percent Similarity:	90.000	Percent Identity:	80.000	
alignment_block:				
US-09-444-281-36 x MMU08210	..			
Align seg 1/1	to: MMU08210	from: 1	to: 2651	
3 ArgTTPProTIRPProTrrpararglys 12				
2025 ACGTGACCTTGCTGGCCCTGGAGGTCCTCGG 2054	:::			
seq_name: gb_ro:AF289665				
seq_documentation_block:				
LOCUS AF289665 107257 bp DNA ROD 14-AUG-2000				
DEFINITION Mus musculus EIF4H gene, partial cds; LIMK1 gene, complete cds; and				
EUN gene, partial cds.				
ACCESSION AF289665				
VERSION AF289665.1	GI:9800517			
KEYWORDS				
SOURCE	house mouse.			
ORGANISM	Mus musculus			
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
AUTHORS	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.			
TITLE	1 (bases 1 to 107257)			
JOURNAL	Direct Submission			
	Submitted (26-JUL-2000) Genome Technology Branch, National Human			
	Genome Research Institute, 49 Convent Dr. Rm. 2N02, Bethesda, MD			
	20892, USA			
FEATURES	location/Qualifiers			
source	1..107257			
	/organism="Mus musculus"			
	/strain="129/Sv"			
	/db_xref="taxon:10090"			
	/chromosome="5"			
	/clone="4.2j20"			
mRNA	/clone_lib="Research Genetics CTRB-CJ7-B BAC library"			
	complement(<7676..7763)			
	/product="EIF4H"			
	complement(<7676..7734)			
CDS	/codon_start=1			
	/product="EIF4H"			
	/protein_id="AAF9335.1"			
	/db_xref="GI:9800519"			
	/translation="MADPFTYDDRAVSFGGGRG"			
mRNA	complement(join(24480..25812,25920..26077,26336..26391,			
	26274..28430,30220..30285,30530..30589,32612..32743,			
	33634..33720,34257..34440,37341..37507,37617..37722,			
	38745..38951,41098..41207,41439..41577,55071..55167,			
	56168..57026)"			
	/product="LIMK1"			
CDS	complement(join(25650..25812,25920..26077,26336..26391,			

US-09-444-281-36 x AP003754/rev ..

Align seg 1/1 to reverse of: AP003754 from: 1 to: 129052

3 ArgTrpProTrrPrrProTrrPrrArgArgLys 12

53243 CGCTGGCCTTGTCGCTGACGCGGG 53214

seq_name: gb_htg:AC091250

seq_documentation_block:
LOCUS AC091250 200849 bp DNA HTG 11-APR-2001
DEFINITION Mus musculus chromosome 5 clone RP23-315E2 strain C57BL6/J, WORKING
DRAFT SEQUENCE, 7 unordered pieces.
AC091250
AC091250.1 GI:13592171
HTG: HTGS_PHASE1, HTGS_DRAFT.
KEYWORDS house mouse.
SOURCE
ORGANISM

REFERENCE
AUTHORS
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 200849)

Ayala, K., Beckstrom-Sternberg, S.M., Benjamin, B., Blakesley, R.W.,
Bouffard, G.G., Brinkley, C., Brooks, S., Dietrich, N.L., Granite, S.,
Guan, X., Gupta, J., Ho, S.-L., Idol, J.R., Karlins, E., Lee-Lin, S.-Q.,
Legaspi, R., Lim, M., Maduro, Q.L., Maduro, Y.B., Masello, C.,
Mastrian, S.D., McCloskey, J.C., McDowell, J., Pearson, R., Prasad, A.,
Shevchenko, Y., Snyder, B., Stanlipop, S., Thomas, J.W., Thomas, P.D.,
Tjongson, E.E., Touchman, J.W., Tsugeon, C., Vogt, J.L., Walker, M.A.,
Weatherly, K.D., Zhang, L.-H. and Green, E.D.

NIH Comparative Sequencing Initiative
Unpublished
2 (bases 1 to 200849)

Green, E.D.
Direct Submission
Submitted (11-APR-2001) NIH Intramural Sequencing Center, 8717
Grovermont Circle, Gaithersburg, MD 20877, USA

Genome Center
Center: NIH Intramural Sequencing Center

Center code: NISC
Web site: http://www.nisc.nih.gov
Contact: nisc_mouse@nih.gov
Project Information

Center project name: aty
Center clone name: 315E02

Summary Statistics

Sequencing vector: plasmid, n/a; 100% of reads
Chemistry: Dye-terminator Big Dye 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 197727 bases at least Q40
Consensus quality: 198423 bases at least Q30
Consensus quality: 198823 bases at least Q20
Insert size: 20200; agarose-ftp
Insert size: 200249; sum-of-ctrls
Quality coverage: 9.87x in Q20 bases; agarose-ftp
Quality coverage: 9.95x in Q20 bases; sum-of-ctrls

* NOTE: This is a 'working draft' sequence. It currently
* consists of 7 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 2812: contig of 2812 bp in length
* 2813 2912: gap of unknown length
* 2913 9301: contig of 6389 bp in length
* 9302 9401: gap of unknown length
* 9402 17173: contig of 7772 bp in length
* 17174 17273: gap of unknown length
* 17274 26044: contig of 8771 bp in length
* 26045 26144: gap of unknown length

FEATURES
source
1..200849
/organism="Mus musculus"
/strain="C57BL6/J"
/db_xref="taxon:10090"
/chromosome="5"
/clone="RP23-315E2"
/clone_1id="RPCI mouse BAC library 23"
1..2812
/note="assembly_fragment"
2913..9301
/note="assembly_fragment"
9402..17173
/note="assembly_fragment"
17274..26044
/note="assembly_fragment"
26145..65929
/note="assembly_fragment"
vector_end:sp6
vector_end:right
66030..109036
/note="assembly_fragment"
109137..200849
/note="assembly_fragment"
clone_end:r7
vector_end:right

BASE COUNT 53444 a 49164 c 49019 g 48614 t 608 others
ORIGIN

alignment_scores:
Quality: 69.00 Length: 10
Ratio: 7.667 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:
US-09-444-281-36 x AC091250 ..

Align seg 1/1 to: AC091250 from: 1 to: 200849

3 ArgTrpProTrrPrrProTrrPrrArgArgLys 12

148453 AGTGGCCTTGTCGCTGACGCTCGG 148482

seq_name: gb_ro:AF210429

seq_documentation_block:
LOCUS AF210429 456 bp mRNA ROD 30-APR-2001
DEFINITION Mus musculus group X secretory phospholipase A2 (Plazg10) mRNA,
complete cds.
AC091250
AF210429.1 GI:12003290

KEYWORDS
SOURCE
ORGANISM

house mouse.
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 456)

REFERENCE
AUTHORS
Morikawa, Y., Saiga, A., Yokota, Y., Suzuki, N., Ikeda, M., Ono, T.,
Nakano, K., Fujii, N., Ishizaki, J., Arita, H. and Hamsaki, K.
Mouse group X secretory phospholipase A2 induces a potent release
of arachidonic acid from spleen cells and acts as a ligand for the
phospholipase A2 receptor
Arch Biochem Biophys. 381 (1), 31-42 (2000)

JOURNAL
MEDLINE
PUBMED
REFERENCE
AUTHORS
20470496
11019817
2 (bases 1 to 456)
Morikawa, Y., Saiga, A., Yokota, Y., Suzuki, N., Ikeda, M., Ono, T.,

TITLE
 JOURNAL
 FEATURES
 source
 gene
 CDS
 BASE COUNT
 ORIGIN
 alignment_scores:
 Quality: 67.00 Length: 10
 Ratio: 7.444 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 80.000
 alignment_block:
 US-09-444-281-36 x AF210429
 Align seg 1/1 to: AF210429 from: 1 to: 456
 1 IIELEAUGTPTPTPTPTPTPTPTPTPTAAG 10
 156 CTTGTTATTGTGTGGCTTGGTGGCCATGGAGA 185
 seq_name: gb_ro:AF166097
 seq_documentation_block:
 LOCUS AF166097 1040 bp mRNA ROD 06-DEC-1999
 DEFINITION Mus musculus group X secreted phospholipase A2 (Pla2g10) mRNA,
 complete cds.
 ACCESSION AF166097
 VERSION AF166097.2 GI:6525307
 KEYWORDS
 SOURCE
 ORGANISM
 Mus musculus.
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 1040)
 2 (bases 1 to 1040)
 3 (bases 1 to 1040)
 Valentin,E., Ghomashchi,F., Gelb,M.H., Lazdunski,M. and Lambeau,G.
 On the diversity of secreted phospholipases A(2). Cloning, tissue
 distribution, and functional expression of two novel mouse group II
 enzymes
 J. Biol. Chem. 274 (44), 31195-31202 (1999)
 200002639
 Erratum: [[published erratum appears in J Biol Chem 2000 Jan
 21;275(3):2246]]
 2 (bases 1 to 1040)
 Valentin,E., Ghomashchi,F., Gelb,M.H., Lazdunski,M. and Lambeau,G.
 Direct Submission
 Submitted (07-JUL-1999) IPMC, CNRS, 660, route des Lucioles,
 Valbonne 06560, France
 3 (bases 1 to 1040)
 Valentin,E., Ghomashchi,F., Gelb,M.H., Lazdunski,M. and Lambeau,G.
 Direct Submission
 Submitted (06-DEC-1999) IPMC, CNRS, 660, route des Lucioles,
 Valbonne 06560, France
 Sequence update by submitter
 REMARK

```

COMMENT      On Dec 6, 1999 this sequence version replaced gi:1614695.
FEATURES
  Source      location/qualifiers
              1..1040
              /organism="Mus musculus"
              /db_xref="taxon:10090"
              /chromosome="16"
              1..1040
              /gene="Pl2g10"
              /gene="Pl2g10"
              /EC_number="3.1.1.4"
              /note="Ca2+-dependent secreted phospholipase A2"
              /codon_start=1
              /product="group X secreted phospholipase A2"
              /protein_id="AA04498.2"
              /db_xref="GI:525308"
              /translation="MLLLILLDGGPGESEARRSHVYKRGELLEAGTLDDCVGPSS
              PMAWMYGCVGLGSGEPPRAIDMCCYHHDCYSRAODAGCSPKLDRYPKCKMDHHTI
              LCGAENKCOQLRCDEBLAYCLAGETVHLKTYLFFPSILCEKDPKCN"

BASE COUNT   244 a      258 c      277 g      261 t

ORIGIN
alignment_scores:
  Quality:    67.00      Length:    10
  Ratio:      7.444      Gaps:    0
Percent Similarity: 90.000      Percent Identity: 80.000

alignment block:
US-09-444-281-36 x AF166097 ..

Align seg 1/1 to: AF166097 from: 1 to: 1040

1 ILeLeuAgtTPrProTrrPrProTrrParg 10
      ::::| | | | | | | | | | | | | |
330 CTGTTATGTGGCCTTGTCGCCATGAGACA 359

seq_name: gb_pl:ATPPHYB
seq_documentation_block:
LOCUS      ATPHYB      3850 bp      mRNA      PLN      13-SEP-1994
DEFINITION Arabidopsis thaliana phyB mRNA for phytochrome.
ACCESSION  X17342
VERSION    X17342.1 GI:16422
KEYWORDS   photoreceptor; phytochrome.
SOURCE     thale cress.
ORGANISM   Arabidopsis thaliana
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
            Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
            1 (bases 1 to 3850)
            Sharrock, R.
            Direct Submission
            Submitted (21-DEC-1989) Sharrock, R., Montana State University,
            Department of Biology, Bozeman, MT 59717 USA
            2 (bases 1 to 3850)
            Sharrock, R.A. and Quail, P.H.
            Novel phytochrome sequences in Arabidopsis thaliana: structure,
            evolution, and differential expression of a plant regulatory
            photoreceptor family
            Genes Dev. 3 (11), 1745-1757 (1989)
            90108670

Location/Qualifiers
1..3850
/organism="Arabidopsis thaliana"
/strain="Columbia"
/db_xref="taxon:3702"
/dev_stage="3 week old green tissue"
/tissue_type="leaf"
/clone_lib="lambda gt10"
/clone="A7-5"
93..3611
/note="phyB photoreceptor"

```

```

/codon_start=1
/protein_id="CAA35222.1"
/db_xref="GI:16423"
/db_xref="SWISS-PROT:P14713"
/translation="MWSVGSGSGGKGGKGGRCGEFEFSSSHTPNNRRGGEDQASSGKKS
LPRSSNTEMSKMAIQQYIVDARLAAVEEGSGESGEKFEVDVSDQKTTTGVSPEDQIT
ATLSRIORGQYIQPFQCMIAVDSSSFRIIGYSNNAEMGLIMQSVPTLEKPEILLAG
TQVRSITFSSSSSTILBRFAVAREITLTLNFWHISKNTGKFFVALIHRIDGVVILPEY
AREDEPALSTAGAVOSOKLAVRAISQLOAGGDIKLCTVVEVSVDLGVDRVMY
KRHEDEGVYAESKRKRDLEPYGLCHYPATDIOASPELEKONRVMTVDCNATPVLY
VODRITQSKCLVGSITLRRAGHGSQYMANMSIASLMAVITINGNDDGSNVAAGRS
SMWLQVLCVCHTSSRCIPPLRYACFEFLMQAGLOIMELIOLAIMSEKRYLRTOTL
LCMDLRLDSPAGIVTQSPSITMDLVKDCGAFLYHGKTPUGVAPSEVQIKDYEMILA
NHADSDSLTSDSLGDAGYPGAAALGDVAGCMAVAYITKRFPLWFSRHTAKETKMGGA
KHHPEDKDGQRMHPRSSFEQFLAEVYKSRQSPWETEMDAIHSIQLILRDSFESEA
MKNVYDVGVQPCRMAGEQIDELGFAVAREMYLLETAIVPIFVADAGCINGMWNK
LALTEGLSVEAMGKSLVSDLIYKRNATVTKLSTRALRDEDEKNVYKLTFSPELO
GKAVFVYVNAACSSKDIYLNITVGYCPGQDPTSOXITWMDKFINQGDYKALVHSPNELL
PLIFADENTCCLEMMAMKELTIGMSRSEYITKMTIYEVGSCOMLKGDPALRKEMIV
LHNAIQGQDIDKPPPEFDNRNGKFVQALITANKVSLDGAVITGAFCLQIPSELOQA
LVAORRDECTCFKAKELAYTCQVINKMPLSGMRPANSLLBATDINEOKOLLETFSVC
EKOSIRIVGMDLESIEDGSEYFLGVSINAVISOAMPLILRRGQILINDIPE
EIKSIEYGRQIRITQOLLAFFELLSITPAQSOEWVEYTHLSQSKOMDGFPAITERR
MACPGECILPEPELYRDMHRSRMTSPBGLISVCKLIKIMNGEYQYIRESRSYFLIITII
LELPEPRKRPLSTRSGSDMLMMPY"
BASE COUNT      1021 a      734 c      1013 g      1082 t
ORIGIN
      polyA_site
      /note="polyA site"
      3850
alignment_scores:
      quality:      64.50      length:      11
      ratio:      6.450      gaps:      1
Percent Similarity:      90.909      Percent Identity:      81.818
alignment_block:
US-09-444-281-36 x ATPHYB      ..
Align seg 1/1 to: ATPHYB from: 1 to: 3850
      3 ARGTPPProTrp...TTPProTrpArgArgLys 12
      |||||
      119 CGGTGGCGGTGGCGGTGGCGGTGGCGGAGAGA 151
seq_name: gb_pl:ATPHPTOCHB
seq_documentation_block:
LOCUS      ATPHPTOCHB      6509 bp      DNA      PLN      14-JUN-1993
DEFINITION      Arabidopsis thaliana phytochrome b gene, exons 2 and 3.
ACCESSION      L09262.1
VERSION      GI:166793
KEYWORDS      photoreceptor; phytochrome B.
SOURCE      Arabidopsis thaliana (strain Landsberg erecta) DNA.
ORGANISM      Arabidopsis thaliana.
Eukaryota: Viridiplantae: Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta: Magnoliophyta: eudicotyledons: core eudicots;
Rosidae; eustoids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE
AUTHORS      Reed,J.W., Nagpal,P., Poole,D.S., Putruya,M. and Chory,J.
TITLE      Mutations in the gene for the red/far red light receptor
phytochrome B alter cell elongation and physiological responses
throughout Arabidopsis development
JOURNAL      Plant Cell 5, 147-157 (1993)
MEDLINE      93200802
FEATURES
      source
      Location/Qualifiers
          1..6509
          /organism="Arabidopsis thaliana"
          /strain="Landsberg erecta"
          /db_xref="taxon:3702"
          2201..2298
          4463..4450
          /number=1

```

```

exon                               4551..5358      /number=2
intron                             5359..5448
exon                               5449..5739
intron                             5740..6115
                                     /number=3
3'UTR                             6369..6509
BASE COUNT      1823 a      1192 c      1459 g      2035 t
ORIGIN

alignment_scores:
  quality:      64.50      Length:      11
  Ratio:        6.450      Gaps:      1
Percent Similarity: 90.909      Percent Identity: 81.818

alignment block:
US-09-444-281-36 x ATHPHTOCHB ..

Align seg 1/1 to: ATHPHTOCHB from: 1 to: 6509

3 ArgTTPPrPrPr...TTPPrTTPArGArGlys 12
|||||
2325 CGGTGCGCGGTGGCGGTGGCGGTGGCGAGAGA 2357

seq_name: qb_pl1:AC005724

seq_documentation_block:
LOCUS      AC005724      86671 bp      DNA      PLN      05-APR-2000
DEFINITION      Arabidopsis thaliana chromosome II section 109 of 255 of the
complete sequence. Sequence from clones MSF3.
ACCESSION      AC005724 AB002093
VERSION
KEYWORDS
SOURCE
ORGANISM
Arabidopsis thaliana
Thale cress.
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1 (bases 1 to 86671)
Lin,X., Kaul,S., Rounsley,S.D., Shea,T.P., Benito,M.-I., Town,C.D.,
Feild,C.Y., Mason,T.M., Bowman,C.L., Barnstead,M.E.,
Feldhym,T.V., Buell,C.R., Ketchum,K.A., Lee,J.J., Ronning,C.M.,
Koo,H., Moffat,K.S., Cronin,L.A., Shen,M., VanAken,S.E., Umayam,L.,
Rallou,L.J., Gill,J.E., Adams,M.D., Carrera,A.J., Creasy,T.H.,
Goodman,H.M., Somerville,C.R., Coppenhaver,G.P., Preuss,D.,
Niernan,W.C., White,O., Eisen,J.A., Salzberg,S.L., Fraser,C.M. and
Venter,J.C.
Sequence and analysis of chromosome 2 of the plant Arabidopsis
thaliana
Nature 402 (6763), 761-768 (1999)
JOURNAL
MEDLINE      20083487
PUBMED      10617197
2 (bases 1 to 86671)
Lin,X.
Direct Submission
Submitted (09-MAR-2000) The Institute for Genomic Research, 9712
Medical Center Dr., Rockville, MD 20850, USA
On Dec 17, 1999 this sequence version replaced gi:4185128.
The sequence and annotation of chromosome 2 were merged from those
of the individual clones on this chromosome after removing
overlaps. For detailed information, please see the TIGR web site
(http://www.tigr.org/tdb/at/at.html).

COMMENT
Genes were identified by a combination of three methods: Gene
prediction programs including GRAIL
(http://athurh.epm.ornl.gov/pub/xcgrail), GeneFinder (Phil Green,
University of Washington), Genscan (Chris Burge,
http://genomic.stanford.edu/GENSCAN.html), and NetPlantane
(http://www.cds.dtu.dk/services/NetGene2/), searches of the
complete sequence against a peptide database and plant EST
databases at TIGR, and manual curations based on those analyses.

```

Annotated genes are named to indicate the level of evidence for their annotation. Genes with similarity to other proteins are named after the database hits. Genes without significant peptide similarity but with EST similarity are named as 'unknown' proteins. Genes without protein or EST similarity, that are predicted by two or more gene prediction programs over most of their length are annotated as 'hypothetical' proteins. Genes encoding tRNAs are predicted by tRNAscan-SE (Sean Eddy, <http://genome.wustl.edu/eddy/tRNAscan-se/>). Simple repeats were identified by RepeatMasker (Arian Smil, <http://ftp.genome.washington.edu/RM/RepeatMasker.html>). Genes are numbered from the top to bottom of the chromosome.

We thank the CSHL/WashU/ABI consortium for sequencing BAC clones F6p23, F5j6, T17A5, and T13L16, the ESSA group for sequencing clone F13D4, and Scott Jackson, Jiming Jiang, Klaus Meyer, Eric Richards and Satoshi Tabata for helpful assistance. In addition, we would like to thank the TIGR Bioinformatics Department, especially Lixin Zhou, Hanif Khalak, Michael E. Heaney, Lily Fu, Feng Liang, Jeremy Peterson, Michael Holmes, and Delwood Richardson for software and database support.

This work was supported by the National Science Foundation, Department of Energy and the US Department of Agriculture.

Address all correspondence to: at@tigr.org.

FEATURES
source

misc_feature

repeat_region

mRNA

gene

CDS

mRNA
gene
CDS

```
1. 86671
/location/Qualifiers
/organism="Arabidopsis thaliana"
/cultivar="Columbia"
/db_xref="taxon:3702"
/chromosome="II"
1. >86671
/note="Sequence from clone MSF3"
/complement(345..409)
/rpt_family="(GAAA)n"
/join(<1030..2192,2354..2494,2618..2896)
/gene="At2g18630"
<1030..2896
/gene="At2g18630"
/note="MSF3.1: unusual splice site at second intron; GA
instead of conserved GT at donor site"
/join(1030..2192,2354..2369)
/gene="At2g18630"
/note="unknown protein"
/codon_start=1
/protein_id="AAD08932.1"
/db_xref="GI:4185129"
/translation="MGKSSSKSKNVFEGSFSTFVQIKINSEYTEHLSYERACSEDP
KLESFDSALHRTNRVINKLASGVEIKSLSPDSLRVYQCLDMNODVKKVILDDKED
IWNODLESLVNLFEETAKTMDCELENCINARSOVITIOFAVNOFEENEDKEN
RKYKTELELKRKFVAGPEPTKEFFALDYKQVMMLEHLKRLDKRLNIKT
MRRYSNMVFAVAFSVLIJSVAAVAAPVVAALAGALAVPGSVGKWCMTMYKKE
KVYAGKEITTSIRIKITYSIKENDNISILRKAVELESILKAEKRIITEKEKRLA
IDETKKLDVFTETIEELGEHAGKYCSDVTAKRVILQRIIRYPAGSPKDEAPWTEW
"
/complement(3613..4761)
/gene="At2g18640"
/complement(3613..4761)
/gene="At2g18640"
/note="MSF3.2: contains GB:122347"
/complement(3627..4745)
/gene="At2g18640"
/codon_start=1
/product="putative geranylgeranyl pyrophosphate synthase"
/protein_id="AAD08933.1"
/db_xref="GI:4185130"
/translation="MEAGNIFLYLLVFLSLHEVFTTLKGLSPANTRRLLRLHIPI
KSPVAALFAKRDREFLDSSIKLVNEEDDFGSEDFEYPMISKAETINRLDLAIPL
IEPINIHAKRYAILLAGKRVPLILACELGSEERLAIQACAVAMHITMSLID
DLPGMNDLDRGKPTTHKVFGESEVAILSGALLALAEHETLEADVSKKMWRAVEL
AKSIGTKLIVAGQAKDLSSGELDNDVGLDELEYTHAKTCSLLSEASVATVAGGCT
EKETEKVNFARCIIGLLFQVVDLIDLETKSSSEELGKTAGKDVAGKLTYPVIGVEKS
```

```
KEFVEKLKRDAREHLGFDSDKVKPLIALTNFIANRNH"
complement(<6733..>8004)
/gene="At2g18650"
complement(<6733..>8004)
/gene="At2g18650"
/note="MSF3.3"
/complement(6733..8004)
/gene="At2g18650"
/codon_start=1
/product="putative C3HC4-type RING zinc finger protein"
/protein_id="AAD08934.1"
/db_xref="GI:4185131"
/translation="MNRILPEMSTQNLSSPPPLPLKSTNSLSNLSKTPNLL
LIILILIFISGLIHILVFLTPSPRESREDFEDVWTALOGLOOLFNLDHSGVDO
SLIDTLPVFNKYSIVGLKISPDPCVLCIEETDKRLPLKCSHAFHVECDITWLS
HSQELPRLSNLSGSESHNNSSYLVLVEESGRMVAVPLENSOLGVDVNDSES
TRISGRSCDPOGMDGLDEKYEPLVGLKSPFNIDHVEGSGOKNSISGNKAND
GRCLSMGSTEYIMDQENTLKVHYSTKLSGKDRVPSHRYVMSCEGDFYKIGEKV
VERSEFSLKTIWLGKREKOKGTARSDCSFVSSSLRPFNHRIPEESLSNSES
LETPTSPFARTMHMLAGRONKIYQSTSNV"
9406..9449
/rpt_family="(GA)n"
10570..10636
/rpt_family="(TA)n"
complement(join(<11029..11303,11403..>11508))
/gene="At2g18660"
complement(<11029..>11508)
/gene="At2g18660"
/note="MSF3.4: predicted by genescan"
complement(join(11029..11303,11403..11508))
/gene="At2g18660"
/note="hypothetical protein"
/codon_start=1
/protein_id="AAD08935.1"
/db_xref="GI:4185132"
/translation="MAVKEVVMVIVPAQIILAPIEAAGKAVYDPPYTSACYGTOR
ETLIVGVKNNLMONGRACGRYRVRCIGATYFNDRACGTGRVDVKKVDFCPCNGDL
NLSRDAFRIANTDAGNIRVYVTP"
11334..11401
/rpt_family="(TA)n"
<13342..>13887
/gene="At2g18670"
<13342..>13887
/gene="At2g18670"
/note="MSF3.5"
13342..13887
/gene="At2g18670"
/codon_start=1
/product="putative C3HC4-type RING zinc finger protein"
/protein_id="AAD08936.1"
/db_xref="GI:4185133"
/translation="MPTNNYRISGEPPSTPSHPPEKPKRIISLFLVGYIMSEFP
LFLYLIGASLIILPLLSSLRHRRHRRRRRRROBSSGLSSRPVKLPQKRSPEST
YTRYESDVCVDFGEGOGWCNRLPGGHVFRKCVDRVDTLLKASTCPICRAKRVLMEE
DPQGEIWMRCGHRSSLLDL"
14514..15636
/gene="At2g18680"
14514..15636
/gene="At2g18680"
/note="MSF3.6"
14653..15516
/gene="At2g18680"
/note="unknown protein"
/codon_start=1
/protein_id="AAD08937.1"
/db_xref="GI:4185134"
/translation="MFSVLPPLLNCLVLYFNALIKPEITNLLESSLLPMDPNT
PEFAHLMRVAVDFRQVNSLYIFIAVSSINLSTLVMAVASLTHRDSPEIKDPF
ILTLKYMGPPLVNTYIYVLSLQWLPFLFYLSIVFSTSLDLSLAASKRLFTYFAV
FESYLAIWNLSNYSIILEDYVIGQALGAKAKIYKGMKPKFLNLPGLISPGIYOI
LRIVDMSSSPSVTLTGTGLVMSVYVVRMPOLVYTYAIVPCKSLQODDSLIDVEYTK
LSSTTLMGGLP"
17474..18643
```



```

GEODIGPPPGPGCIGYPCGMAGPKGEMGPRGYKGMVSTGACPPCEGRCPPGEAG
EKGDVSGOARGGPGOGITGPKITGPPCIGDKDGPGLIPKMGSGAGVGRGSPGHOIL
AGVPGPGCTGKGPGDKGPGGCGGLPGVSGPPKGEPPGRTIGPGOSIMOKGDGGR
GPVGQRPGRPGRCQPKGEQGPPIPGPGLGPGIKGDKSPKRTGPRGCVGPPVAGLGG
EKKEKQSGSEPPGLKGQGVKGETGYPGPSGDIGAPGVQGYPLPGRGLVDKVPQ
PGKQGVVGRASDQHIYDVVLKMIQEOLEAVASAKREALGAGMWGLPDPGPPGYP
GKQGNHGPGRGIPGIVGAVQIGNTPKKGKEDRGEMGHGPMGPICIDLP
PGRPGQAINGKDDGRSGAPGEGRGRGPVGLPFCPEPACTLGASAVTSARLTFR
GSTKGP"
exon      4311..4385
           /gene="alpha2 (IX) collagen"
           /label=ex2
           /number=2
           /evidence=experimental
           5646..5681
           /gene="alpha2 (IX) collagen"
           /label=ex3
           /number=3
           /evidence=experimental
           5768..5830
           /gene="alpha2 (IX) collagen"
           /label=ex4
           /number=4
           /evidence=experimental
           7048..7101
           /gene="alpha2 (IX) collagen"
           /label=ex5
           /number=5
           /evidence=experimental
           7214..7249
           /gene="alpha2 (IX) collagen"
           /label=ex6
           /number=6
           /evidence=experimental
           7347..7370
           /gene="alpha2 (IX) collagen"
           /label=ex7
           /number=7
           /evidence=experimental
           7630..7683
           /gene="alpha2 (IX) collagen"
           /label=ex8
           /number=8
           /evidence=experimental
           8010..8063
           /gene="alpha2 (IX) collagen"
           /label=ex9
           /number=9
           /evidence=experimental
           8168..8215
           /gene="alpha2 (IX) collagen"
           /label=ex10
           /number=10
           /evidence=experimental
           8344..8400
           /gene="alpha2 (IX) collagen"
           /label=ex11
           /number=11
           /evidence=experimental
           8485..8538
           /gene="alpha2 (IX) collagen"
           /label=ex12
           /number=12
           /evidence=experimental
           8828..8881
           /gene="alpha2 (IX) collagen"
           /label=ex13
           /number=13
           /evidence=experimental
           9268..9321
           /gene="alpha2 (IX) collagen"
           /label=ex14
           /number=14

exon      9413..9466
           /evidence=experimental
           /gene="alpha2 (IX) collagen"
           /label=ex15
           /number=15
           /evidence=experimental
           9572..9625
           /gene="alpha2 (IX) collagen"
           /label=ex16
           /number=16
           /evidence=experimental
           12979..13032
           /gene="alpha2 (IX) collagen"
           /label=ex17
           /number=17
           /evidence=experimental
           13393..13446
           /gene="alpha2 (IX) collagen"
           /label=ex18
           /number=18
           /evidence=experimental
           13672..13725
           /gene="alpha2 (IX) collagen"
           /label=ex19
           /number=19
           /evidence=experimental
           14530..14574
           /gene="alpha2 (IX) collagen"
           /label=ex20
           /number=20
           /evidence=experimental
           14967..15020
           /gene="alpha2 (IX) collagen"
           /label=ex21
           /number=21
           /evidence=experimental
           15284..15337
           /gene="alpha2 (IX) collagen"
           /label=ex22
           /number=22
           /evidence=experimental
           15543..15596
           /gene="alpha2 (IX) collagen"
           /label=ex23
           /number=23
           /evidence=experimental
           15674..15745
           /gene="alpha2 (IX) collagen"
           /label=ex24
           /number=24
           /evidence=experimental
           15857..15932
           /gene="alpha2 (IX) collagen"
           /label=ex25
           /number=25
           /evidence=experimental
           16069..16113
           /gene="alpha2 (IX) collagen"
           /label=ex26
           /number=26
           /evidence=experimental
           16207..16239

exon      64.00      Length: 9
           Quality:      Gaps: 0
           Ratio: 8.000      Percent Identity: 77.778
Percent Similarity: 88.889

alignment block:
US-09-444-281-36 x MMA2IXCOA/rev ..
Align seg 1/1 to reverse of: MMA2IXCOA from: 1 to: 19479
```

4 TrpProTrrPrrPrrPrrPrrPrrPrr 12
|||||
4355 TGGCCTGTGGTGGCTGGAGACCGG 4329

seq_name: gb_hc9:AL356097_0

seq_documentation_block:

WPCOMMENT

Sequence split into 4 fragments LOCUS AL356097 Accession AL356097

Fragment Name

AL356097_0

AL356097_1

AL356097_2

AL356097_3

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

1 (bases 1 to 355026)

Submitted (07-APR-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk
requests: clonerequests@sanger.ac.uk
On May 15, 2001 this sequence version replaced gi:8077023
gi:9797422.

----- Genome Center

Center: Sanger Centre

Center code: SC

Web site: http://www.sanger.ac.uk

Contact: humquerry@sanger.ac.uk

----- Project Information

Center project name: bAI80A14

----- Summary Statistics

Assembly program: XGAP4; version 4.5

Sequencing vector: M13; M7815; 43% of reads

Sequencing vector: plasmid; L08752; 56% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Consensus quality: 33302 bases at least Q40

Consensus quality: 346727 bases at least Q20

Insert size: 351326; sum-of-coverage

Insert size: 176896; 2.8% error; agarose-fp

Quality coverage: 3.95x in Q20 bases; sum-of-coverage

Quality coverage: 8.06x in Q20 bases; agarose-fp

Draft Sequence Produced by Whitehead Institute/MIT Center for

Genome Research, 320 Charles Street,

Cambridge, MA 02141, USA

http://www-seq.wi.mit.edu.

* NOTE: This is a 'working draft' sequence. It currently

* consists of 38 contigs. The true order of the pieces

* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will

* be preserved.

1 16446: contig of 16446 bp in length

16447 16546: gap of 100 bp

16547 22746: contig of 6200 bp in length

22747 22846: gap of 100 bp

22847 45482: contig of 22636 bp in length

45483 45582: gap of 100 bp

45583 49251: contig of 3663 bp in length

49252 49351: gap of 100 bp

49352 53239: contig of 3888 bp in length

FEATURES

Source

* 53240 53339: gap of 100 bp
* 53340 58178: contig of 4839 bp in length
* 58179 58278: gap of 100 bp
* 58279 71754: contig of 13476 bp in length
* 71755 71854: gap of 100 bp
* 71855 76641: contig of 4787 bp in length
* 76642 76741: gap of 100 bp
* 76742 80415: contig of 3674 bp in length
* 80416 80515: gap of 100 bp
* 80516 89902: contig of 9387 bp in length
* 89903 90002: gap of 100 bp
* 90003 93661: contig of 3659 bp in length
* 93662 93761: gap of 100 bp
* 93762 101785: contig of 8024 bp in length
* 101786 101885: gap of 100 bp
* 101886 109333: contig of 7448 bp in length
* 109334 109433: gap of 100 bp
* 109434 121288: contig of 11855 bp in length
* 121289 121388: gap of 100 bp
* 121389 128767: contig of 7379 bp in length
* 128768 128867: gap of 100 bp
* 128868 131917: contig of 3050 bp in length
* 131918 132017: gap of 100 bp
* 132018 138795: contig of 6778 bp in length
* 138796 138895: gap of 100 bp
* 138896 144660: contig of 5765 bp in length
* 144661 144760: gap of 100 bp
* 144761 156587: contig of 11827 bp in length
* 156588 156687: gap of 100 bp
* 156689 158939: contig of 2252 bp in length
* 158940 159039: gap of 100 bp
* 159040 161353: contig of 2314 bp in length
* 161354 161453: gap of 100 bp
* 161454 165170: contig of 3717 bp in length
* 165171 165270: gap of 100 bp
* 165271 168479: contig of 3209 bp in length
* 168480 168579: gap of 100 bp
* 168580 172188: contig of 3609 bp in length
* 172189 172288: gap of 100 bp
* 172289 177084: contig of 4796 bp in length
* 177085 177184: gap of 100 bp
* 177185 199876: contig of 22692 bp in length
* 199877 199976: gap of 100 bp
* 199977 222186: contig of 22210 bp in length
* 222187 222286: gap of 100 bp
* 222287 240235: contig of 17949 bp in length
* 240236 240335: gap of 100 bp
* 240336 263715: contig of 23380 bp in length
* 263716 263815: gap of 100 bp
* 263816 269270: contig of 5455 bp in length
* 269271 269370: gap of 100 bp
* 269371 271745: contig of 2375 bp in length
* 271746 271845: gap of 100 bp
* 271846 277024: contig of 5179 bp in length
* 277025 277124: gap of 100 bp
* 277125 280037: contig of 2913 bp in length
* 280038 280137: gap of 100 bp
* 280138 307328: contig of 27191 bp in length
* 307329 307428: gap of 100 bp
* 307429 313985: contig of 6557 bp in length
* 313986 314085: gap of 100 bp
* 314086 330085: contig of 16000 bp in length
* 330086 330185: gap of 100 bp
* 330186 334887: contig of 4702 bp in length
* 334888 334987: gap of 100 bp
* 334988 355026: contig of 20039 bp in length.

Location/Qualifiers

1. 355026

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="1"

/clone="RP11-180A14"

/clone_1lb="RPCT-11.1"

LTR
/note="3' LTR"
join(23418..23748,25558..25589,25715..26480,26838..26965)
/note="hypothetical protein"
/codon_start=1
/protein_id="BAA92399.1"
/db_xref="GI:7228439"
/translation="MGANCCIAKKEPPQCVPIEVSARFNRHSESMFSRMDNRTHI
EDIMPPALFENHSGSIPKESGSIAPTDGFSNGSPSDMKNLAKHSDKRRSS
KIARSDRGSTNSSPKESGSIAPTDGFSNGSPSDMKNLAKHSDKRRSS
RHSGLPTDADSMKARRSPGYOLRYQVSDSKLPSLNEGASPEGRPSMNLVYCN
DISAVSGHSSSDGMSMRTFSEVVASORERMSVDELIGSVSKMTRSNANPNTH
SPDOEYCKTCLIKLKERSTWNOELAVAVILGHHYHADCDSIADKDKDPPEV
CHGEKCYKILGKLESKTKNNIPKMYIVDVNLDGSKHOKKEIPLLEELYSOARSM
RELKSRIGANDMTLNDLRLEGFS"
complement(join(27953..28624,29565..29918,30369..30646,
31026..31491))
/note="GSGS C96615(C10106),C26336(C12127),D21959(C10106)
correspond to a region of the predicted gene.
Similar to NAM (A021889)"
/codon_start=1
/protein_id="BAA92400.1"
/db_xref="GI:7228440"
/translation="MESTRDVLPFGCFHFKPDTLISHYLLKIKHGIKEVEIPEV
DLYHEPMDLPKADVPYQDNKMHFAARDKYPNGSRNRATVAGVWSTKDAIK
MKQITGPKTKLVEHGRPPGRTKRTMHEHYIDREQACPKDKDAIVLCRITRN
DMIPNGNELNDSHPPEYDAPPSYISTEQLNPAEPVGAELVAAVTAEPDGVTS
ATTANIPSPSDINLDDWLNLELDFEPEOSLASADLSPDONVSSWGLAKVE
QYSSPNNVVDITEXILPEEDVNIILHPTGDDFNMLONLDOYPIAYADWVGLOKE
ELMSPOANAEPSQSNMEADNGIIRIRSKMTPETSPQKGTOKAKMVGIKMATS
SSSINOTIKFENGSLVHEKQKADADVASTKRSADGKSTELSSNRGFLGRIRNAFAC
CSDAWMNLVAGPAVAVVLAHIGRLGLSQROOH"
complement(38747..38998)
/note="hypothetical protein"
/codon_start=1
/protein_id="BAA92401.1"
/db_xref="GI:7228441"
/translation="MGHFLYLAADTARPNMMAVGPPEVRYHVRHGTPEVPLVLCRPD
DNGPCRAHMAKMLGLLVLHFGCCDDEPHHPSGTSI"
complement(39315..39726)
/note="3' LTR"
complement(join(40209..40396,40917..43545))
/note="EST C28952(C62945) corresponds to a region of the
predicted gene.
Similar to maize transposon MudR mudra protein isolog.
(AC003981)"
/codon_start=1
/protein_id="BAA92402.1"
/db_xref="GI:7228442"
/translation="MSKYTPQIVHGEENIRGPGVDLSDPVMTSKGIDRAEPTQ
STYWLKGRIDQEVITMSVSVSRATEGFWELMPDSTDAMRYEAFEMSWP
LVIYVSEKIDINVMQTEDVGPINADIVVPSQNEQNEQREBOAMGADGRRV
LIVDEEDSDNEEDDASDEEDGADINAMADESGVLSIGDHPMYKENEV
IEGARVYHDEKFAVKHMAVSLQREFRVAESTYNTVEKCECPMVAVYKGMN
DYKSYIVTEHKYGVGEVYKIRNTISAFMASEMSSVYVNGCEPKSIRIIRTEKKF
VTISAKMRAKOKITTEKRYGTFEASYNLPLAIIAQRNNNTYYDLHTFTSVDR

LTR
/note="3' LTR"
join(44572..44723,44846..44938,46366..46378)
/note="hypothetical protein"
/codon_start=1
/protein_id="BAA92403.1"
/db_xref="GI:7228443"
/translation="MALPGRVSPKPMMAAASASPPQPRGHMLVLTGTFGRVYSQ
RLIAGWRSVSGTSPAKKTELEMGDASIDPATSSRCV"
complement(join(48119..48257,48429..48703,49975..50139))
/note="hypothetical protein"
/codon_start=1
/protein_id="BAA92404.1"
/db_xref="GI:7228444"
/translation="MALVNDKDEDDVNAACARRAPLVIGAPMGFVLQARLIGEC
EPRDRARMAITISPCALSLFSKMTYIYVILCKRYRPLIYTKESKPKCRIT
ASLARHGELEICLARASSALDRVRYKILCTDFVLMLVKRCIYINPHIRTH
GSMVSTLPDAPVYSTRTELAGIYDTCIGS"
complement(join(51918..52637,52684..53366))
/note="Similar to Arabidopsis thaliana chromosome 1 YAC
YUP8H12R sequence, C2-HC type zinc finger protein
C.e-WY11. (AC002986)"
/codon_start=1
/protein_id="BAA92405.1"
/db_xref="GI:7228445"
/translation="MASCINGEELHVRVYHRLVYASDESIRPHVLAWSNDLIRPTI
QVSMFCIYKPGSTGDFHDVVAFAAGLPSLNLHFPPLAGRIVSNPCSGLEPIHCNQ
GAELVAGADVALASIDGTGASVAGKILLPAGCVVALSVGVSPGAGGFVAMCTNH
VYVDSGSLMLVANSIELARSGTLAAGRPNDHRSFPPSPSYGASIDEAFPLDG
ARQADIALRFEORATRTVOAVATLWMLAAVGSRDARCMVWVWGRRRLTSS
PELRAMSVYCNVNTTPVAAVETMEIORKPLAIVASAROMAIPAAYGHROPEVDV
VEHNAKGRQYIDTASVGLSGPVAVYTAFASEVYTDTHGHANALPTSSSARLCT
GVQVLAARSGGDSWIASALMLPLAALLESDEBLGKIFPVYAEVYLGRPASSSA
KRAGMTSKG"
complement(57570..58009)
/note="3' LTR"
complement(join(58018..58891,58980..62098))
/note="Similar to Arabidopsis thaliana chromosome II BAC
F9013 genomic sequence, putative retroelement pol
polyprotein. (AC006248)"
/codon_start=1
/protein_id="BAA92406.1"
/db_xref="GI:7228446"
/translation="MRVNLQAGLMTAIDPGYAEPRDMAALSAILQAVPRBMLGLA
KHDTAKAMDAIKTRVGVDRYREAKKEGFRQFMSMKKEEPPEEFAMRTAVAD
IRDMGVKAMDAIKTRVGVDRYREAKKEGFRQFMSMKKEEPPEEFAMRTAVAD
DEEGSDGKILYITEOMQARVYKORQESQNSGRAPGQVQAAKEEPTLLMAHYG
ATGANSRDISRVCNFCNDEPGHVAROORCKPGRORGBANVYQAAKEEPTLLMAHYG
VSLAGEATLGRTPGGOEVHLTEKYYIIDLHDEGGEFEVYRMDPDIQGANHMTGVASAP
AELDTCVGYTKVFGGSVIEILOGKTYVFRCKNGDRSLDYITPKLRKNIISVGR

alignment_scores:

Quality:	Length:
Ratio: 8.000	9
Percent Similarity: 88.889	Gaps: 0
Percent Identity: 88.889	

alignment_block:

US-09-444-281-36 x AP001366

Align seg 1/1 to: AP001366 from: 1 to: 146081

3 ArgTTProTTPtrTPtrTPatgag 11
|||||

Mon, Jan 7 10:42:19 2002

74276 CGTGGCCATGCTGCGTTGGAGAGG 74302

us-09-444-281-36.rge

Page 13

THIS PAGE BLANK (USPTO)